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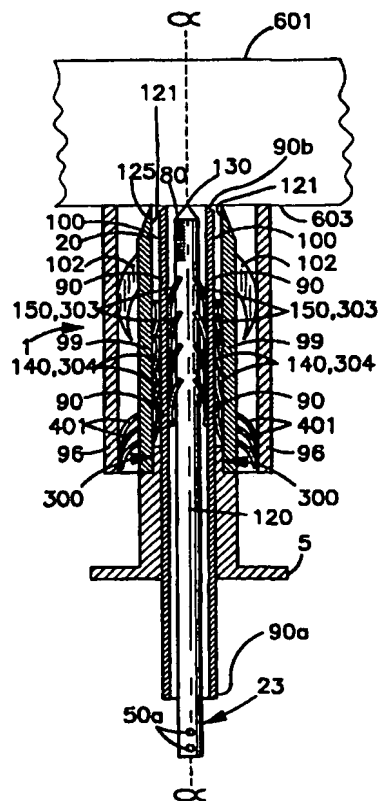
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**(54) Title:** A TISSUE ANCHOR SYSTEM**(57) Abstract**

A tissue anchoring system, including a tissue anchoring device (1), and tissue anchors (20). The tissue anchoring device (1) includes a housing (2), and a tissue anchor (12) positioned on the tissue anchoring device (1) that will be inserted into a tissue or secured onto a tissue. The device (1) may optionally have a plunger assembly (5) slidably positioned in the housing to assist advancing anchor into a tissue. The tissue anchor (20) has a barb end (21), and a shaft (120). The barb end (21) is adapted to resist removal from a tissue after the anchor has been inserted. The tissue anchor shaft (120), and the barb end (21) may be hollow. The anchor shaft (120) has an attachment member (23) distal from the barb end (21) for direct attachment (23) to a tissue or for attachment of sutures or slings. The tissue anchors (20) may include an adjustment mechanism for intra- or post-operative adjustments.



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## A Tissue Anchor System

### Technical Field

This invention relates to a tissue anchor and applicator for supporting a suture, sling member, or other device for use in a surgical procedure. In particular, this invention relates to  
5 tissue anchors for use in surgical treatment of urinary incontinence.

### Background Art

One problem this invention intends to address is urinary incontinence, but the invention is applicable to a broader range of problems. The background, however, will discuss urinary incontinence as an example.

10 Urinary incontinence is an involuntary release of urine when increases in abdominal pressure, caused by sneezing, coughing, or exercising, are not uniformly transmitted to the proximal urethra, resulting in urine "leakage." Moderate stress urinary incontinence ("SUI") is inconvenient and can be a social and hygienic problem, while severe SUI can be disabling. SUI occurs in women and is caused by either hypermobility of the bladder neck and proximal urethra  
15 (excessive downward and rotational movement of the bladder neck) or intrinsic sphincter deficiency.

Several defects can result in loss of support of the bladder neck. Examples of these defects are: (a) breakage or abnormal stretching of the passive supports of the bladder and urethra (those connective tissues supporting these structures); or (b) loss of the active support of the  
20 bladder neck, vagina, and rectum provided by the levator ani muscle. For instance, one example of passive support failure is paravaginal "defects" caused by separation of the vaginal wall from the pelvis caused by breakage or stretching of connective tissue (pelvic organ prolapse). Such separation results in downward rotation of the vaginal wall, which in turn results in downward rotation of the bladder neck because the bladder is partially supported by the vagina. An example  
25 of loss of active support is loss of muscle tone in the levator ani muscle. This muscle operates as a floor, or platform, supporting the bladder during standing and normal activity. Normal muscle tone of the levator ani relieves the passive support of the pressures caused by overlaid abdominal viscera. Loss of muscle tone may result in downward rotation of the bladder.

Treatment of SUI caused by hypermobility of the bladder and proximal urethra requires  
30 supporting the bladder neck, generally at the urethrovesical junction ("UVJ"). Correction requires support of the UVJ area, which helps balance increased abdominal pressures and which allows

the bladder neck to properly compress and close in response to increased pressures, thus preventing urine leakage. Support may be provided in various surgical procedures, including anterior colporrhaphy, retropubic urethropexy, vaginal needle urethropexy, and suburethral sling procedures. One particular type of surgery is briefly discussed—retropubic urethropexy (“RU”). In RU, a series of sutures are used to support the UVJ. The ends of the sutures attach at various points in the body cavity depending upon the type of failure (loss of active support, failure of a portion of passive connective tissue, etc.). The sutures support and properly elevate the bladder neck at the UVJ to maintain the compressibility and pliability of the urethra and to avoid compromising the urethral sphincteric mechanism.

In various RU procedures, different anchoring tissues are used to support the sutures. These anchoring tissues include pubocervical fascia, pubourethral ligaments, Cooper’s ligaments, rectus fascia, and inferior rami of the pubic bone. One RU procedure, generally known as the Marshall-Marchetti-Kantz procedure, uses several sutures, with one end of each suture attached to the vaginal wall adjacent to the urethra, straddling the UVJ bilaterally. The opposite suture end passes through the retropubic periosteum.

In another RU procedure, known as the Burch procedure, the first end of a series of sutures is positioned in both sides of the vagina wall (straddling the urethra) below Cooper’s Ligament 602, and the other end of the sutures is positioned in Cooper’s Ligament 602, thereby supporting the vaginal wall and the UVJ.

RU procedures generally support an area by using sutures, and additional support is provided by using slings (either man-made materials or tissue grafted material) placed under the area to be supported and sutured into an anchoring tissue, such as in suburethral sling procedures (e.g. a Goebell-Stoeckel procedure).

Problems associated with surgical correction of the failed support mechanisms include under- or over-correction of the UVJ. The surgeon must determine the degree of support necessary to properly elevate and support the UVJ to properly address the SUI problem. This determination must be made both pre- and intra-operatively. Too little elevation causes SUI to remain, although the degree of SUI may be reduced. Too much elevation can result in voiding dysfunction (reduced capacity or inability to void), prolonged catheterization, and the need for postoperative correction.

The incidence of postoperative urinary retention can be as high as 30% at two weeks after

surgery, and 5% of patients have postoperative urinary retention that persists. Many patients with less severe cases of postoperative obstructive symptoms also benefit greatly. Symptomatic detrusor instability represents the bladder's response to increased outlet resistance caused by an improperly tensioned sling. The incidence of postoperative irritative symptoms secondary to detrusor instability can be as high as 20%. Appropriate tensioning of the sling minimizes persistent incontinence and voiding dysfunction.

Generally, all of the above procedures require placement of sutures into an anchoring tissue or tissues and may also require placement of sutures into supporting devices, such as slings. Placement is provided through suturing needles, either straight or arcuate, or the use of needle suturing devices, such as push and catch systems, or non-adjustable fascial attachment systems. The suture placement in the tissue can be completely through the tissue or partially through the tissue.

#### **Summary of the Invention**

Accordingly, the invention is a tissue-anchoring system, including a tissue-anchoring device and tissue anchors. The tissue-anchoring device includes a housing and a tissue anchor. The tissue anchor is positioned on the tissue-anchoring device that is advanced into a tissue. The device may optionally have a plunger assembly slidably positioned in the housing to assist advancing the anchor into a tissue. The tissue anchor has a barb end and a shaft. The barb end is adapted to resist removal from a tissue after it is inserted. The tissue anchor shaft and the barb end may be hollow. The shaft of the anchor has an attachment member distal from the barb end so that the attachment member may attach directly to tissue or attach to tissue using sutures or a sling. The tissue anchors may be adjustable.

#### **Disclosure of the Invention**

It is an object of the invention to provide a tissue-anchoring device that does not require suturing to attach to tissue.

It is an object of the invention to provide a tissue-anchoring device that may be repeatedly reloaded with anchors.

It is an object of the invention to provide a tissue-anchoring device that may insert completely through a tissue or partially through a tissue.

It is an object of the invention to provide an alternative to sutures in relatively inaccessible areas.

It is an object of the invention to provide a tissue-anchoring device that can be hand-held or used with endoscopic surgery techniques and devices, especially flexible endoscopes.

It is an object of the invention to provide an adjustable tissue-anchoring device to adjust the placement of devices or tissues attached to the anchor postoperatively without further  
5 invasive surgery.

It is an object of the invention to provide an adjustable tissue-anchor device attachable to tissue using sutures or staples when a surgeon deems it more suitable.

### Description of the Drawings

Figure 1 illustrates an embodiment of a tissue-anchor delivery device.

10 Figure 1a illustrates another embodiment of a tissue-anchor delivery device.

Figure 2 illustrates another embodiment of a tissue-anchor delivery device having a plunger with the delivery device shown as a cross-section.

Figure 3a illustrates a thimble-like embodiment of a tissue-anchor delivery device.

15 Figure 3b illustrates another embodiment of a tissue-anchor delivery device that may be used in connection with an endoscope wherein the delivery device is shown in cross-section to illustrate the plunger positioned therein.

Figure 3c illustrates the configuration of the tip used in connection with the tissue-anchor delivery device shown in Figure 3b.

20 Figure 4a illustrates an anchor having a ringed attachment member and finger projections for a barb.

Figure 4b illustrates an anchor having an umbrella-like barb.

Figure 4c illustrates an anchor having an attachment member a disk barb and an attachment member comprising holes.

Figure 4d illustrates an anchor having an arrow shaped barb and finger-like barbs.

25 Figure 4e illustrates an anchor having a T-shaped barb having holes in the attachment member.

Figure 4f illustrates an anchor having a star-shaped barb.

Figure 4g illustrates the anchor shown in Figure 4b having a barb retaining device in the form of a collar positioned over the umbrella-shaped barb.

30 Figure 4h illustrates another barb retaining device, namely a sleeve positioned over the barb.

Figure 5a illustrates an embodiment of an anchor having a ratcheting device configured therein as a groove.

Figure 5b illustrates an embodiment of an anchor having a shaft positioned therein, wherein ratcheting devices are positioned on only one side of the anchor and a corresponding side of the shaft.

Figure 5c illustrates the embodiment of the anchor shown in Figure 5a shown with a shaft inserted therethrough, illustrating first and second ratcheting devices engaged.

Figure 6a illustrates an embodiment of an attachment member having holes extending therethrough.

Figure 6b illustrates another embodiment of an attachment member wherein holes are configured with a tab near the bottom of the shaft.

Figure 6c illustrates another embodiment incorporating a ring-shaped attachment member.

Figure 6d illustrates another embodiment of an attachment member in the form of a clamp.

Figure 6e illustrates another embodiment of an attachment member comprising finger-like projections and holes.

Figure 6f illustrates another embodiment of an attachment member having annular projections positioned on the shaft.

Figure 6g illustrates another embodiment of an attachment member having indentations configured in the shaft.

Figure 7a illustrates an embodiment of the delivery device having a plunger and using a T-shaped anchor.

Figure 7b illustrates another embodiment of the delivery device having a plunger with a disc-like end designed to engage the shaft of the anchor and anchor head.

Figure 8a illustrates an embodiment of an adjustable tissue-anchoring device.

Figure 8b illustrates another embodiment of an adjustable tissue-anchoring device.

Figure 8c illustrates another embodiment of an anchor body with the barbs removed and having annular projections thereon to engage a retaining device.

Figure 9a illustrates an embodiment of a tissue-anchor retaining device shown as a disk.

Figure 9b illustrates another embodiment of a tissue-anchor retaining device shown as

disk having suture apertures therein.

Figure 10a illustrates an embodiment of a shaft having annular projections as ratcheting devices wherein the attachment member is rotatably attached.

5 Figure 10b illustrates an embodiment of a shaft similar to the embodiment shown in Figure 10a wherein the ratcheting devices are finger-like projections.

Figure 11a illustrates an embodiment of a thimble-shaped delivery device wherein sutures are attached near the barb.

Figure 11b illustrates another embodiment of a delivery device.

10 Figure 11c illustrates another embodiment of a delivery device having a pen-shaped housing.

Figure 12 illustrates an embodiment of a delivery device wherein the anchor slides onto the housing and the shaft slides through the housing and the anchor.

Figure 13 illustrates an embodiment of a delivery device comprising two plungers.

Figure 14 illustrates an embodiment of an adjuster cylinder.

15 Figure 15 illustrates an exploded view of an embodiment of the invention incorporating an adjuster cylinder.

Figures 16a-16g illustrate the steps of placement of an adjustable anchor and shaft within a tissue. In each Figure 16a-16h, the plunger, anchor body, the sleeve, and the adjuster cylinder are shown as cross-sections. The shaft and interleaving members are shown in full view against  
20 those cross-sections.

Figure 17 illustrates a cross-sectional view of an embodiment of an adjustable anchor using threads to allow adjustment of the shaft wherein the shaft is shown in full view.

Figure 18a illustrates an adjustable anchor attached to Cooper's Ligament 602.

Figure 18b illustrates a close-up view of the embodiment in Figure 18a.

25 Figure 18c illustrates a close-up view of the embodiment in Figure 18a wherein corresponding threads are configured on the shaft and housing to allow the shaft to be adjustable.

Figure 18d illustrates an adjustable anchor having a body that accommodates multiple shafts.

30 Figure 19a illustrates a cross-sectional view of another embodiment of the invention wherein the shaft and attachment member are not connected, and wherein the ratcheting device comprises corresponding threads positioned on the housing and on the shaft. Figure 19a also



illustrates the attachment member as rotatable about the housing. The shaft is shown in full view, and an attachment member is shown operatively attached to the housing.

Figure 19b illustrates an embodiment similar to that shown in Figure 19a wherein the ratcheting device on the shaft comprises annular projections. The shaft is shown in full view.

5 Figure 20 illustrates a schematic view of an anchor deployed within a flexible delivery device that also incorporates a micro-video camera and video screen.

### **Best Mode for Carrying Out the Invention**

Illustrations of construction, design, and methods of operation of the invention are set forth below with specific references to the Figures. However, it is not the intention of the  
10 inventor that the scope of his invention be limited to these embodiments.

Figure 1 illustrates one embodiment of the tissue-anchor delivery device 1. Delivery device 1 comprises a housing 2 and one or more finger grips 3, which comprise flats 4 on opposing sides of housing 2 designed to enable a surgeon to hold device 1. Finger grips 3 may be rings 6 (see Figure 3b) through which the surgeon inserts fingers, a thimble (see Figure 3a),  
15 a handgrip, or any other shape that enables a surgeon to hold delivery device 1. Finger grips 3 are unnecessary if housing 2 is adapted to be hand-held, such as a pen-shaped housing 11 shown in Figure 1a.

Figures 1a and 3a illustrate an anchor stay 50. Anchor stay 50 comprises a portion of housing 2 adapted to retain an anchor 20 (not shown) thereto, as described below. In the  
20 embodiment shown in Figure 3a, anchor 20 (not shown) will slip over anchor stay 50, while in the embodiment shown in Figure 1, anchor 20 will clip onto stay 50 using clip member 12.

Anchor stay 50 is a rigid projection extending away from the plane of the finger grip flats 4. Anchor stay 50 may have a clip member 12 positioned near tip 57 to removably couple anchor 20 to stay 50. Clip member 12 may include a compression-fit C-type ring for gripping the shaft  
25 of an anchor 20. Anchor stay tip 57 may also be designed to reversibly mate with an anchor 20 that is positioned on or against tip 57.

Alternatively, as shown in Figure 2, delivery device 1 may include a plunger 5 slidably positioned in hollow barrel portion 7 of housing 2 and operated by a surgeon. Plunger 5 has an anchor end 10 adapted to engage an anchor 20. Anchor end 10 may be a flat end to push anchor  
30 20, a pointed end adapted to engage a hollow shaft portion of an anchor 20 (see Figures 3a and 11c), a clip member 12 (see Figure 1) positioned on end 10 designed to couple to anchor 20.

Though not shown, clip member 12 may alternatively be positioned on anchor shaft 120 (seen in Figures 4a-4h). Anchor end 10 can have any structure engaging anchor 20 that allows plunger 5 to advance anchor 20 into a tissue and uncouple therefrom, leaving anchor 20 in tissue.

Other embodiments of delivery device 1 are shown in Figures 3a and 3b. These  
5       embodiments also comprise a housing 2 and finger grip(s) 3. As shown in Figure 3a, housing 2 and finger grips 3 comprise a thimble 3a, while Figure 3b shows finger grips 3 as rings 6. Viewing Figure 3b, housing 2 has a flexible hollow barrel portion 7. A flexible plunger 5a is slidably positioned in hollow barrel portion 7, which comprises a sleeve within which flexible plunger 5a slides. This embodiment is suitable where the invention will be used in conjunction  
10       with an endoscope or in anatomic areas that are hard to reach manually with a rigid device. Hollow barrel portion 7 may be rigid, but is preferable flexible so that it may be used in conjunction with an endoscope. The hollow barrel portion 7 length may range from 1/4-inch to one or more feet, depending upon the particular application, and the length of flexible plunger 5a is constructed accordingly.

15       Viewing Figure 3b, hollow barrel portion 7 has a detachable tip 8. Tip 8 screws onto hollow barrel portion 7 using corresponding threads 8b, 8c which allows a surgeon to change tips 8. Viewing Figure 3c, tip 8 has a knife-edge 13 to penetrate tissue. Tip 8 may have a slot 9 starting at the edge 8a of tip 8 and extending down tip 8, substantially along the longitudinal axis  $\alpha$  shown in Figure 3c.

20       Viewing Figure 1a, delivery device 1 may also comprise a stop 60, shown as an outwardly projecting ridge 60a, that is shown located on the external walls 7a of the hollow barrel portion 7 in Figure 1a. Ridge 60a controls the extent of insertion of hollow barrel portion 7. Alternatively, viewing Figure 1, finger grips 3 can act as stop 60. The need for a separate delivery device 1 may be eliminated if anchor 20, later described, is itself constructed to guide  
25       the placement of anchor 20 into tissue by hand.

As shown in Figure 4a-f, anchor 20 can have a wide variety of shapes depending upon the application. Anchor 20 has a barb end 21 and a shaft 120 and may also comprise an attachment member 23 (see Figures 4a, 4c, and 4e) positioned on shaft 120 distal from barb end 21. Inclusion of shaft 120 is optional. Please note that in later embodiments, the anchor shaft is  
30       also referred to as shaft 120 (e.g. Figures 16a-16g). Viewing Figure 11a, if anchor 20 lacks shaft 120, attachment member 23, later described, should be placed on barb end 21 (not shown), or the

sutures 450 should directly attach to barb 102. Shaft 120 may be from 1/4 inch to a foot or more in length, depending upon the application.

Viewing Figure 4a, barb end 21 is adapted to resist removal of anchor 20 after anchor 20 has been inserted in or through a tissue and anchored by barb 40. For example, barb 40 may comprise: a series of downwardly-pointing fingers 30 (also referred to as wing-like projections 30) (see Figure 4a); a downwardly-pointing conical umbrella 31 (see Figure 4b); a flat area 32, such as a disk 32a (see Figure 4c) or an arrow end 32b (see Figure 4d); a T-shaped head 33 (see Figure 4e); a star-shaped head 34 (see Figure 4f); or any other head shape that will resist removal of anchor 20 from tissue. Barb 40 may be flexible or collapsible, such as the umbrella 31 shown in Figure 4b. Preferably, umbrella 31 is collapsible.

Viewing Figures 4g and 4h, when barbs 40 are collapsible (generally, foldable inward toward anchor shaft 120), a barb retaining device 94, such as a ring 95 (see Figure 4g), or a sleeve 96 (see Figure 4h and Figure 15), may slide over shaft 120 to collapse barbs 40. Retaining device 94 is retractable down shaft 120, and may be placed over a collapsed barb 40 to hold barb 40 in a collapsed position. When barb end 21 is inserted into tissue, barb-retaining device 94 move down shaft 120 as barb 40 slides through barb-retaining device 94 and deploys into the tissue. Barb 40 stays collapsed, however, as long as the barb sidewalls contact tissue. After penetration of tissue, barb 40 expands, and resists removal from the tissue. Barb 40 also creates resistance by the counter-traction that occurs when barb 40 is imbedded within the tissue.

Materials of construction for anchor 20 are preferably biologically-inert plastics, thin stainless steel, or other non-reactive materials that can co-exist within a tissue with little or no adverse patient reaction. Portions of anchors 20, however, should be sufficiently rigid to insure that anchor 20 cleanly and precisely penetrates the tissue at the desired location and that a deployed barb 40 will resist removal from tissue. Generally, the portion of shaft 120 attached to plunger 5 or attached to anchor stay 50 should be fairly rigid.

Barb end 21 may also be shaped to penetrate a tissue, such as is shown in Figures 4a, 4b, 4f, and 4h. When anchor 20 has a barb end 21 that is ill-suited to penetrate a tissue (see Figures 4c and 4e), anchor 20 is preferably used with delivery device 1 having hollow barrel portion 7 that is shaped to penetrate a tissue. For example, see sharp end 125a shown in Figure 7a.

Viewing Figure 7a, if barb 40 comprises a T-shaped head 33, anchor 20 may be loaded into delivery device 1 using forceps. Shaft 120 is a flexible material that allows placement of T-

shaped head 33 as shown. Sharp end 125a of hollow barrel portion 7 is adapted to penetrate a tissue. A portion of shaft 120 extends through slot 9 configured in housing sidewalls 9a. Preferably, barb end 21 has a larger cross-section than shaft 120, and prevents removal of barb end 21 through slot 9. Plunger 5 engages anchor 20. Though not shown, anchor end 10 of plunger 5 may comprise a clip member 12 designed to partially encircle shaft 120 (see Figure 1a for example).

Viewing Figure 7b, anchor end 10 of plunger 5 may comprise circular-shaped disk 43 with a slot 41 into which shaft 120 is slidably inserted. Shaft 120 passes adjacent to plunger 5 and exits the bottom of delivery device 1, eliminating the need for slot 9. Alternatively, but not shown, anchor end 10 may have slot 41 that works in conjunction with slot 9 located on sidewalls 9a. In such an embodiment, slot 41 aligns with slot 9, enabling shaft 120 to exit hollow barrel portion 7 below anchor end 10.

Viewing Figures 6a-6g, attachment member 23 is a structure generally located on shaft 120 for engaging a material, such as a suture or tissue sample, a second barb end, or a tissue-retaining device 27 (shown in Figures 9a and 9b). Tissue-retaining device 27 may comprise a disk-like washer or button 62 (see Figures 9a and 9b) or simply a lip that operates in conjunction with attachment member 23 and reversibly mates with attachment member 23. Viewing Figures 6a-h, attachment member 23 comprises one or more openings 50a in shaft 120 (or a tabbed portion 23c thereof) (see Figures 6a and 6b); a ring 51 (see Figure 6c), a tissue clamp 52 (see Figure 6d), ratcheting devices, such as series of projection fingers 55 (see Figure 6e) or annular projections 56 (see Figure 6f) extending radially from shaft 120, or a series of indentations 58 in shaft 120 (see Figure 6g), all of which comprise interleaving members.

Alternatively, viewing Figure 6e, attachment member 23 may comprise a combination of the above, such as openings 50 and fingers 55. Viewing Figure 6d, clamp 52 has first and second reversibly interlocking surfaces 53, 54 that may clamp onto tissue. As shown in Figures 6e and 6f, fingers 55 or projections 56 may attach a tissue-retaining device 27 (not shown, see Figure 16f) onto shaft 120, thereby allowing retaining device 27 to compress and retain a tissue between barb end 21 (or tip 130) of the anchor shaft 120 and retaining device 27. This embodiment is well-adapted to support organs whose connective supportive tissues have weakened or failed, such as the transvaginal sacral or sacrospinous colpopexy later described.

Viewing Figures 9a and 9b, tissue-retaining device 27 is an annular disk-like structure,

sometimes referred to herein as a "porcupine" button 62. Button 62 has a center hole 404 sized for positioning around shaft 120 and attachment member 23. The top surface 27a of button 62 may be arcuate-shaped, and have a series of projections 29 extending outwardly from top surface 27a. Projections 29 grasp tissue to prevent button 62 from migrating. Viewing Figure 9b, to resist migration of button 62 relative to tissue, button 62 may have a series of holes 61 extending therethrough around the periphery for suturing button 62 to tissue. Alternatively, button 62 may be constructed of a mesh material, allowing sutures to extend therethrough. A surgeon must ensure contact between button 62 and shaft 120 does not interrupt blood supply to that area in contact with tissue-retaining device 27.

If attachment member 23 is as a ratcheting device, such as fingers 55 projecting from the exterior sidewalls of shaft 120 as shown in Figure 6e, or a series of indentations 58 (see Figure 6g), and tissue-retaining device 27 is button 62, button 62 may slide up or down shaft 120. However, button 62 is retained in a given position on shaft 120 by the inter-mating of the ratcheting devices on shaft 120 and within button center hole 404. Center hole 404 may have a series of fingers 402, or other projections, extending outwardly to interact with the interleaving members or ratcheting devices on shaft 120 to accomplish the same function. Generally, button 62 has a first interleaving member (fingers 402) on the interior walls of hole 404, while anchor 20 will have a second interleaving member (such as projections 701 in Figure 8c) positioned thereon. The first and second interleaving members cooperate to resist movement of button 62 relative to anchor 20. Alternatively, a C-clip or other clip member 12 on delivery device 1 may be used as a retaining device 27 to keep anchor 20 properly positioned relative to the tissue.

Viewing Figures 4a, 4c, and 4e, if sutures are placed through attachment member 23, they are preferably installed when anchor 20 is loaded into delivery device 1 so that: (a) the surgeon does not have to position sutures through openings 50a after anchor 20 is attached, and (b) if the sutures are sufficiently long to reach back to the surgeon during placement of anchor 20, the surgeon will not fish for the sutures in the body cavity. To assist a surgeon in tailoring anchor 20 to a particular application, attachment member 23 may be detachable from anchor 20 (e.g., threaded onto shaft 120, or clipped onto shaft 120 or the base of barb end 21, etc). Shaft 120 has a small cross section at tip 130, much like a needle tip, such as 1-3 mm, to allow easy penetration into tissue.

Viewing Figure 19a, anchor 20 and attachment member 23 may comprise separate

elements. Anchor 20 has a barb 102 positioned on a shaft 120. Attachment member 23 comprises a housing 1000, and shaft 120 is movably inserted through housing 1000. Housing 1000 comprises an upper portion 1005 and a lower portion 1006 rotatably engaged using interlocking lips 1007, 1007a. Interlocking lips 1007, 1007a are slidably engaged, and to assist rotation, a bearing 1009 or other suitable device may be included. Shaft 120 is movably positioned within housing 1000 via corresponding threads 1004, 1004a.

An alternate embodiment is shown in Figure 19b. Housing 1000 and shaft 120 engage each other with interleaving members, or ratcheting devices 140, 150, such as grooves 1001 in housing 1000 and corresponding projections 1001a positioned on shaft 120. Interleaving members 1001, 1001a allow the position of shaft 120 to be fixed relative to housing 1000. Shown on shaft 120 is an attachment member 23 and holes 50a for attaching sutures or other suitable items.

The embodiment shown in Figures 19a and 19b, allows a surgeon to adjust the position of shaft 120 relative to attachment member 23 and items attached to attachment member 23 during placement of the tissue-anchoring system. The embodiment in Figure 19b can be adjusted using an adjuster cylinder 90, and the embodiment in Figure 19a can be adjusted by rotating shaft 120, causing shaft 120 to move up or down.

#### **Examples of Use of the Anchoring System**

Viewing Figure 7a, to set anchor 20 in tissue, a surgeon positions a loaded delivery device 1 against the tissue to be penetrated for insertion of anchor 20. If the delivery device 1 has a hollow barrel portion 7 designed to penetrate tissue, then hollow barrel portion 7 is placed against the tissue, and delivery device 1 is advanced with hollow barrel portion 7 penetrating tissue. When hollow barrel portion 7 has advanced so that sharp end 125a clears the tissue or is embedded therein, plunger 5 is advanced, releasing anchor 20. Hollow barrel portion 7 is withdrawn from the tissue, leaving anchor 20 supported within the tissue or by a tissue surface 601 (not shown). If delivery device 1 has a hollow barrel portion 7 that will not penetrate tissue, then anchor 20 should be equipped with a sharp end 125 (not shown, see Figure 8a) designed to penetrate a tissue. In this embodiment, hollow barrel portion 7 is placed against the tissue, and plunger 5 advanced, forcing anchor 20 into tissue. When anchor 20 has advanced sufficiently into tissue or through tissue, delivery device 1 is withdrawn, leaving barb 40 embedded into tissue, or inserted through tissue.

Viewing Figures 11a-11c, if delivery device 1 has no hollow barrel portion 7, (for example, the anchoring system as shown in Figure 1a) then barb 102 of anchor 20 is adapted to penetrate a tissue. In such an embodiment, anchor 20 is placed against the tissue to be penetrated, the delivery device 1 advanced (or if it is plunger-equipped, plunger 5 is advanced), until anchor 20 has advanced sufficiently into tissue or through tissue, at which time the delivery device 1 is withdrawn.

This anchoring system may also be used in less surgically technical operations for surgeons who are not comfortable with transvaginal approaches in laproscopically-assisted sacral colpopexy operations. The surgeon may place anchor 20 into the anterior longitudinal ligament of the sacrum laproscopically or endoscopically and secure the vaginal vault to the sacrum transvaginally in a laproscopically assisted sacral colpopexy. Alternatively, the system may be used with a microscopic video camera 65 located near tip 8 of delivery device 1 as seen in Figure 20. The anchoring system would be placed into and or through the apex of the vaginal vault using a rigid or flexible delivery device 1, either endoscopically or transvaginally, to fix anchor 20 to suspend the vault to the sacrum. The anchoring system can also be used endoscopically with an operative single port laparoscope thereby obviating the need for multiple puncture sites in the abdomen when performing this procedure. The anchoring system obviates the need for other bulkier surgical instruments currently used for retropubic bladder neck suspension, retropubic Goebell-Stoeckel sling procedures, and multiple abdominal scope port sites. This anchoring system also obviates the need for using bone anchors and obviates the associated well-known complications. In addition, by using the tissue anchoring system and an intervening connector to attach to a sling of harvested tissue, the size of the material harvested or used in the sling can be decreased in size, thereby minimizing the amount of graft tissue or donor tissue needed to be used. In turn, this minimizes the amount of graft tissue and operative harvesting technique, thereby minimizing complications, such as hematoma formation, pain, infection and decreased strength of the upper leg. In addition, the retropubic Goebell-Stoeckel sling could be used using a smaller fascia sling graft or donor graft using an umbrella shaped anchor head 31 with an extended anchor shaft 120 armed at both ends creating a u-shaped anchor that seats into the rectus fascia 603 via an exclusively transvaginal approach.

As indicated, the system can be used with a tissue clamp 52 as attachment member 23 (see Figure 6d). Clamps 52 fix the ends of a fascia sling for the Goebell-Stoeckel fascia sling

procedure and obviate the need for sutures. A general outline of the transvaginal-endoscopic-sacralcolpexy procedure using an endoscope-adapted tissue anchor system follows. The patient is placed into the dorsal lithotomy position and undergoes general or spinal anesthesia. The vaginal cuff is visualized using angled surgical retractors. A transverse incision is made just inside the hymenal ring on the posterior wall and extended down the mid-line of the posterior vaginal wall, sharply and bluntly dissecting the vaginal mucosa from the underlying supportive tissue of the vagina to the apex of the vault. The retroperitoneal space would be accessed using sharp and blunt dissection revealing the sacral promontory to the S2 and S3 areas of the sacrum and the anterior longitudinal spinal ligament.

The vaginal cone speculum is inserted up to the anterior longitudinal spinal ligament. The middle sacral vessels are visualized and isolated to the exterior of the cone using blunt retraction. If this is impossible and there is a significant risk for damage to these vessels, the vessels may be ligated using a hemoclip-extended applicator transvaginally through the vaginal cone superior and inferior to the operative sites. If bleeders are encountered in the retroperitoneal space while dissecting the anterior longitudinal spinal ligament, they could be fulgurated using a Klepinger electrocautery device or hemoclip. The surgeon secures the tip and rim of the vaginal cone against the sacrum maintaining isolation of the operative site while inserting a flexible fiberoptic endoscope into the vaginal cone at the sacrum loaded with the flexible tissue-anchor applicator. The tissue anchor 20 is loaded onto a delivery device 1. The shaft of the anchor is advanced into and parallel to the under surface of the anterior longitudinal spinal ligament for approximately one centimeter. The appropriate angle of insertion is facilitated by the manipulation of the flexible tip of the anchor/anchor applicator and/or endoscope by the surgeon at the operator end of the endoscope. The delivery device 1 is advanced through the operative port of the endoscope thereby advancing the anchor with barb end through the anterior longitudinal spinal ligament to the desired depth. Plunger 5 is depressed and advances anchor 20 into the space anterior to the sacrum but just beneath the anterior longitudinal spinal ligament. Delivery device 1 is withdrawn into the endoscope.

The surgeon visualizes anchor 20 in the anterior longitudinal spinal ligament. The endoscope is withdrawn trailing the sutures which are attached to anchor 20 back into the vaginal cone speculum and towards the surgeon at which time they are grasped in a clamp for later attachment to a synthetic mesh, fascia lata or rectus fascia 603 graft for later suspension of the



vaginal vault to the sacrum. If necessary, several anchors 20 may be placed into the anterior longitudinal spinal ligaments in this fashion.

An extended suction device is used to enhance visualization of the space at the operative site. A suction irrigator device is preferred. The synthetic graft is attached to the trailing sutures from the anchor(s) 20 and then advanced up the vaginal cone speculum, applied to the sacrum and tied in a fashion similar to that done during an endoscopic procedure with an extended knot pusher. This secures the proximal portion of the graft at the anterior longitudinal spinal ligament. The distal portion of the graft is secured to the vaginal vault apex using: (a) the standard transvaginal surgical instruments that one would use during a vaginal hysterectomy, or (b) the adjustable tissue anchor clasp and adjustable suture connector for attachment and adjustment of the vaginal cuff. The graft application is performed entirely in the retroperitoneal space. Before placing the sutures at the vaginal vault apex, the surgeon ensures that the sutures are placed into the graft at a site that allows appropriate length between the vaginal apex and the sacrum minimizing any excessive tension on the mesh or fascial graft. The vaginal cuff is then closed in a manner used in the standard posterior-colporrhaphy procedure, well-described in the gynecologic surgical literature. Because this procedure is carried out endoscopically (or manually) in the retroperitoneal space, there is no need for abdominal entrance and/or closure of the parietal peritoneum over the mesh or fascial graft. This ensures that there are no internal hernias postoperatively, and the operation is totally extra-peritoneal thereby minimizing ileus. The surgeon should be careful and concerned about hemostasis and ensuring that the colon and ureter are not damaged during the procedure. The vaginal cone at its apex should be padded with a soft silicone gasket to ensure that tissues are not unduly traumatized.

Tissue anchor 20 can also be used in the post-operatively adjustable transvaginal-sacrospinous-colpopexy procedure as follows. The patient is placed in a dorsal lithotomy position after undergoing general or spinal anesthesia. This procedure may be performed using local anesthesia for patients who have medical conditions that may be complicated by either spinal or general anesthesia. A weighted speculum is placed in the posterior vault. A Sims retractor is placed anteriorly. The cuff apex is visualized if the uterus and cervix have been removed. In the lateral vaginal fornices, a 0.5-1.0 centimeter (cm) vaginal mucosa incision is made, and a 1-2 cm<sup>2</sup> area of vaginal mucosa undermined at these sites. This reveals the underlying submucosal vaginal-supportive tissue. Tissue-anchor delivery device 1 is placed into the surgeon's hand and

advanced into the vagina. The tissue-anchor delivery device 1 with a loaded tissue anchor 20 is advanced up the vagina and punctures the vaginal tissue through the lateral fornices of the vagina on the patient's right side directly through the area which was previously undermined. Tissue-anchor delivery device 1 places anchor 20 into the sacrospinous ligament as follows. The sacrospinous ligament is located after the surgeon palpates the ischial spine, and anchor 20 is placed approximately 2 cm medial to the ischial spine through the sacrospinous ligament mid-portion. Tissue-anchor 20 deploys into and/or through the sacrospinous ligament, and delivery device 1 is withdrawn from the vagina leaving anchor 20 in position transfixing the vagina vault apex to the sacrospinous ligament. The chosen anchor 20 depends on the surgeon's desires.

Anchor 20 may be adjustable or non-adjustable. Viewing the embodiments shown in Figures 16a-16h or Figure 17, adjustable tissue anchors 20 are used as follows. Anchor 20 is positioned as previously described, transfixing the lateral vaginal cuff apex to the sacrospinous ligament. Delivery device 1 is then removed. This leaves the adjustable shaft 120 in anchor 20 trailing from the lateral vaginal apex into the vaginal canal. A 1-1.5 cm<sup>2</sup> "porcupine" button 62 is then advanced onto shaft 120 and ratcheted into place using the interleaving members 303, 304 or ratcheting devices 140, 150 positioned on shaft 120 and in some instances using the interleaving members (fingers 402) positioned in center hole 404, thereby opposing the lateral vaginal cuff apex to the sacrospinous ligament. The vaginal cuff apex may be clasped or sutured alternatively to anchor 20. The same procedure is carried out on the opposite side. This suspends the vaginal cuff apex to the sacrospinous ligament either unilaterally or bilaterally. The trailing end of shaft 120 may be used for adjusting the vaginal cuff tension postoperatively and then trimmed flush with anchor body 99 when the appropriate tension is achieved. If the surgeon does not want a foreign body in the vagina, a standard adjustable tissue anchor 20 could be used, the vaginal cuff apex sutured to anchor 20, and vaginal suspension tension adjusted postoperatively to ensure patient comfort.

The vaginal mucosa is closed using either running- or interrupted-absorbable sutures over button 62 at the lateral vaginal cuff apices. In addition, button 62 may be sutured to the submucosal vaginal tissue along its circular perimeter before closure of the vaginal mucosa to ensure its appropriate fixation in the tissues. The lateral circular rim of button 62 may be composed of a mesh-like synthetic material, to facilitate penetration of fibroblasts and granulation tissue to ensure fixation of the button in the subvaginal mucosal tissue.

The procedure is preceded by the appropriate preparation of the surgical site with an antiseptic solution such as iodine or other antiseptic and appropriate draping to ensure sterile technique. The patient should also receive pre-operative prophylactic antibiotics, approximately one to two doses, and one to two doses postoperatively after closure of the vaginal mucosa to prevent postoperative infection from a foreign body.

This procedure obviates the need for extensive vaginal dissection as described in other techniques used in vaginal vault suspension such as sacrospinous ligament fixation using the tendon sheath punch, or the hook-like suture carriers. Other vaginal vault suspension procedures require a more extensive dissection of the posterior vaginal vault and submucosal tissue. This more extensive dissection can result in serious hemorrhage. The procedure is done with a minimal amount of vaginal dissection--approximately 4 cm<sup>2</sup> divided at two locations.

There is also minimal manipulation of the sacrospinous ligaments and surrounding tissue. This procedure requires only a single puncture of the sacrospinous ligament. The operative time would be substantially decreased as a result of the simplicity of operative dissection insertion of the tissue anchor system and the ability to adjust vaginal suspension tension postoperatively, as later described. This procedure could be used on an outpatient basis.

#### **Adjustable Tissue Anchor**

An embodiment of an adjustable anchoring system is shown in Figures 8a-8c. Figure 8a illustrates an anchor 20 comprising a body 99 having a hollow chamber 100 therein and a shaft 120 insertable within body 99. Shaft 120 is sized to be at least partially insertable into hollow chamber 100 and moveable relative to hollow chamber 100. Hollow chamber 100 may pass completely or partially through body 99. Viewing Figure 8b, if hollow chamber 100 extends partially through body 99, body 99 preferably has a slot 89 that allows shaft 120 to exit hollow chamber 100 through sidewall 89a. Shaft 120 has a tip 130 adapted to penetrate a tissue. Tip 130 should be fairly rigid to ease tissue penetration, although a flexible tip 130 is preferred for certain procedures, such as transvaginal sacral colpopexy. Viewing Figures 16a-16h and Figure 18b, shaft 120 may have indicia markings 80 thereon (e.g. millimeter (mm) markings or color change markings) as indicators that allow a surgeon to determine the extent of tension adjustments. Shaft 120 may have a cross-section that is round, flat, rectangular, or any other suitable shape.

The end of shaft 120 opposite tip 130 has an attachment member 23 attached thereto or constructed as part thereof. Hollow chamber 100 has a barb end 101 with at least one barb 102

positioned thereon. Barb 102 resists removal of anchor 20 from a tissue after anchor 20 has been inserted into or through tissue. Barb end 101 may be a variety of shapes but is easily insertable into tissue, generally having a sharp end 125 and a collapsible barb 102 for use in conjunction with a barb retainer 94, as previously described.

5 Viewing Figure 8c, body 99 may also have a ratcheting device 701 positioned on body outer walls 703 distal from barb end 101. Ratcheting devices 701 may comprise a series of fingers, annular projections 702, indentations, or threads to adjustably and matingly engage tissue-retaining device 27, such as button 62. Alternatively, tissue-retaining device 27 may comprise a washer-like structure that snaps or clips onto shaft 120 where desired. Though not  
10 shown, body 99 may also have a stop 70 projecting away from body walls on the end distal from barb end 101 to prevent the body 99 from inserting completely through the desired anchoring tissue.

Viewing Figures 5a-5c, the interior walls 121 of hollow chamber 100 may have a first ratcheting device 140. First ratcheting device 140 may be a series of annular ridges, threads,  
15 projections, indentations or other shapes designed to interlock or interleave with a second ratcheting device 150 located on exterior of shaft 120 (such as interlocking male and female threads, indentations and corresponding projections (see Figure 5b), interleaving sidewall fingers 303 shown in Figures 16a-16h, interleaving annular projections 1001a shown in Figure 19b, and Figure 5c). Variation in the number or type of interleaving members or ratcheting devices or their  
20 materials of construction will vary the load-support capability of a particular anchor 20. Ratcheting devices 140, 150 create a two-way ratcheting-type mechanism that positions shaft 120 relative to body 99. Shaft 120 may be adjusted in steps by pulling or pushing shaft 120 relative to body 99 (or turning, in the case of interlocking threads). If ratcheting devices 140, 150 comprise male and female threads, it is desirable that shaft 120 have a ball bearing 15 or other  
25 pivotally rotatable mechanism 14 located in attachment member 23 as shown in Figures 10a and 10b.

Mechanism 14 allows shaft 120 to turn independently of the bottom of shaft 120 where attachment member 23 is located. Mechanism 14 prevents twisting of sutures, slings, or other devices attached to attachment member 23. This twisting may occur during later tension  
30 adjustments, either intra- or post-operative.

Anchor 20 is set by first setting outer hollow chamber 100 and then shaft 120 with the surgeon's hand or with a delivery device 1. A delivery device 1 is preferred because anchor 20 may be relatively small. Viewing Figure 12, delivery device 1 may comprise a housing 2 and finger grips 3, but other previously-described embodiments may also work.

5 Alternatively, anchor delivery device 1 may include a hollow barrel portion 7 wherein at least a portion of shaft 120 is positionable therein. Hollow barrel portion 7 has a lip section or edge 300, sized to fit within hollow chamber 100. Though not shown, a stop member may be positioned on body 99 to prevent body 99 from being deployed completely through tissue. Though not shown, delivery device 1 may optionally include a plunger 5 slidably positioned in  
10 hollow barrel portion 7. Inclusion of a plunger 5 in the embodiment of Figure 12 requires that plunger 5 adapt to allow shaft 120 to pass through or by plunger 5 (e.g., plunger 5 as shown in Figure 7b). In such an embodiment, plunger 5 alone activates shaft 120.

Alternatively, as shown in Figure 13, housing 2 may have a hollow barrel portion 7 that is adapted to have anchor 120 positioned at least partially therein. Hollow barrel portion 7 should  
15 have a sharp end 125a adapted to penetrate a tissue. This embodiment is appropriate when barbs 102 are ill-suited to penetrate a tissue. Delivery device 1 may further comprise one, two or no plungers. In a two-plunger embodiment, a first plunger 301 activates body 99 after hollow barrel portion 7 has penetrated a tissue, and second plunger 302 activates shaft 120 after barbs 102 deploy. Figure 13 illustrates a two-plunger embodiment comprising concentric first and second  
20 plungers 301, 302. A single-plunger embodiment may be used if either plunger 301 or 302 activates either body 99 or shaft 120. However, if a single plunger is adapted to activate shaft 120, the surgeon should rely on the friction between first and second ratcheting devices 140, 150 on body 99 and shaft 120 to cause deployment of body 99 when the plunger is depressed. After barbs 102 deploy, shaft 120 may be accessed through the skin surface 601 (see Figure 16f) by  
25 a surface incision as will be later described. Though not shown, housing 2 may also have a projecting ridge 60a located on the external side of hollow barrel portion 7 to act as a stop 60.

Shaft 120 is adjustable upwardly or downwardly, with or without the aid of adjuster cylinder 90 shown in Figure 14. As used herein, upward and downward shall indicate movement substantially along the central axis  $\alpha$  of anchor 20 or housing 2 as shown in the Figures. Figure  
30 15 illustrates an exploded view of the invention that shows how adjuster cylinder 90 inserts into hollow chamber 100. Viewing Figure 15, adjuster cylinder 90 is a hollow tube having an inner

and outer diameters sized so that the adjuster cylinder 90 may slide over shaft 120 but slide inside of hollow chamber 100, thereby disengaging the interleaving members 303, 304 or ratcheting devices 140, 150. Adjuster cylinder 90 has a lower end 90a, and may have a cutting edge 90b.

Alternatively, the tissue-retaining device may be as shown in Figures 18a-18d. Viewing Figure 18a, pelvis 600 is shown from a retropubic view. Cooper's ligament 602 is located on pelvis 600. Sutured to Cooper's ligament 602 is anchor 20, which has a hollow portion 66 extending therethrough. Shaft 120 inserts through anchor 20 and is shown exiting skin surface 601. Anchor 20 may be configured with one or more suture holes 67 that allow anchor 20 to be attached to Cooper's ligament 602. In this embodiment, anchor 20 lacks a barb 102. Shaft 120 inserts through hollow portion 66. Shaft 120 is adjusted using a shaft 120 having threads 68 (see Figure 18c) corresponding to threads position in the hollow portion 66, or by placing corresponding interleaving members or ratcheting devices 140 within hollow portion 66 and corresponding interleaving members or ratcheting devices 150 on shaft 120. Shaft tip 120b may be sharp so that it can penetrate a tissue and skin surface 601.

Shown in Figure 18b is an enlarged view of Figure 18a, to more clearly show ratcheting devices 150 (interleaving members) on shaft 120. Ratcheting devices 150 (interleaving members) shown in Figure 18b are annular projections similar to those shown in Figures 5c and 8a. Shown in Figure 18c is the same device as shown in Figure 18b, but shaft 120 has threads 68 positioned thereon allowing shaft 120 to rotatably engage corresponding threads on the interior walls of hollow portion 66. Attachment member 23 rotatably attaches to shaft 120 and is shown as a ball and socket 15. Figure 18d illustrates an embodiment wherein anchor 20 has multiple hollow portions 66 so that two shafts 120 can engage anchor 20.

#### **Examples of use of adjustable tissue anchor**

Referring to Figures 16a-16g, one of the procedures where the adjustable anchor is advantageous is the retropubic bladder neck suspension or retropubic Goebell-Stoeckel sling procedure. In this procedure, it is necessary to support the neck of the bladder through the use of a sling. The following application will demonstrate the use of the adjustable tissue anchor in this procedure, using two anchors: one non-adjustable anchor and one adjustable anchor. If the sling is to be tacked to the bladder neck, it is desirable to use two adjustable anchors. The procedure could be performed transvaginally or by laparotomy.

The surgeon first sets a non-adjustable anchor into a suitable connective tissue on one

side of the bladder neck, Cooper's Ligament 602 or rectus fascia using the procedure(s) previously described. Anchor 20 may have a sling attached thereto by sutures. The sling is suspended under the urethra and fixed with sutures. The surgeon attaches a second adjustable anchor to the other end of the sling (by use of sutures or by use of a tissue clamp 52 as attachment member 23 on shaft 120) and loads the adjustable anchor 20 onto delivery device 1. Alternatively, the adjustable tissue anchor 20 shown in Figures 18a-18c could be sutured to Cooper's Ligament 602, rectus fascia 603, or tendon bilaterally and subsequently adjusted postoperatively. This is accomplished using ratcheting devices 140, 150 ratcheted or thread and screw adjustment mechanism (i.e., interleaving members). The surgeon proceeds to set adjustable anchor 20 into a suitable location on the other side of the bladder neck using a delivery device 1 as shown in Figure 12 or in Figures 5a-5c.

Viewing Figures 16a-16g, delivery device 1, with the tissue anchor 20 located therewithin or thereupon, is positioned next to the area where attachment is desired. Delivery device 1 is then advanced forcing shaft tip 130, and the sharp end 125 of body 99 to penetrate into tissue. The barb 102 is retracted or collapsed by barb retainer 94. Retainer 94 remains stationary as the surgeon pushes body 99 into the tissue, and barb 102 moves into or through the tissue via the opening created by body 99. Body 99 and shaft 120 then penetrate and insert through the rectus fascia 603 (tendon or other desired tissue), until barbs 102 are located in the soft subcutaneous tissue between the rectus fascias and the skin surface 601 and deploys. Shaft 120 trails from the tissue into which anchor 20 is embedded. At this point, delivery device 1 may be removed (if desired), and anchor 20 may be anchored at the inferior border of the rectus fascia 603 or underneath the rectus fascia 603 in a sandwich-like fashion by applying a tissue-retaining device 27, such as button 62.

Next, shaft 120 is advanced by hand or with a hemostat, or using adjuster cylinder 90 until shaft 120 contacts the underside of skin surface 601. Adjuster cylinder 90 may comprise a cutting edge 90b for tissue penetration as shown in Figures 16a-16c. For instance, the adjuster cylinder 90 could be advanced over shaft 120 from below and completely through delivery device 1 until adjuster cylinder 90 advances into and punctures skin 601.

Alternatively, if shaft 120 is adapted to penetrate a tissue, shaft 120 may be advanced through skin surface 601 by moving shaft 120 upward or moving the abdominal wall downward, forcing shaft 120 through skin surface 601.

In another method, the surgeon locates the exit point of shaft 120 by pressing the abdominal wall until it contacts shaft 120 and makes a small incision with a scalpel at the exit point. By further depression of the abdominal wall, the surgeon causes shaft 120 to exit through the abdominal wall incision. The surgeon grasps and holds the protruding shaft 120 and removes pressure on the tissue and abdominal wall, thus fully deploying adjustable anchor 20. If this particular procedure is followed, shaft tip 130 need not be adapted to penetrate skin surface 601. Delivery device 1 and anchor 20 shown in Figure 12 or 15 may deploy in this fashion.

Yet another method would be applicable with delivery device 1 having a plunger 5 adapted to activate shaft 120. In such an embodiment, shaft 120 is advanced by operation of plunger 5.

After shaft 120 passes through skin surface 601, it is held above skin surface 601, and the tension on the sutures or sling is adjusted by pulling or pushing shaft 120, using adjuster cylinder 90 to disengage the interleaving members, if needed. Once proper tension is achieved, the adjuster cylinder 90 is removed, and the interleaving members/ratcheting devices 140, 150 engage, fixing shaft 120 in place. Shaft 120 is left trailing above skin surface 601 for further postoperative adjustment.

Viewing Figure 16a, to ease movement of shaft 120 through hollow chamber 100, ratcheting devices 140, 150 (interleaving member 303, 304) are disengaged using adjuster cylinder 90 inserted into hollow chamber 100 between shaft 120 and interior walls 121 (see Figure 5a) of body 99. Alternatively, when shaft 120 first protrudes from the incision, adjuster cylinder 90 may be positioned through the abdominal wall (see Figure 16g). When anchor 20 fully deploys, adjuster cylinder 90 is removed, and shaft 120 is left trailing through skin surface 601 as previously described.

An alternative procedure using a plunger-adapted delivery device 1 follows, using a two-plunger delivery device 1 shown in Figure 13. Initially, delivery device 1 is positioned adjacent to the area to be anchored and housing 2 and hollow body portion 7 pushed upward penetrating the tissue (note the sharp end 125a of hollow body portion 7 may be adapted to penetrate a tissue). Once sharp end 125a is properly positioned for deployment of anchor 20, (generally above the surface of the penetrated tissue), the surgeon will deploy at least first plunger 301 to activate body 99, although it may be desirable to operate both plungers 301, 302. This deploys barb 102 in the soft subcutaneous tissue, superior or exterior to the rectus fascia 603 or other



tissue involved. Shaft 120 is advanced by plunger 302, until shaft 120 penetrates skin surface 601. The surgeon may assist by pressing downward on the abdominal wall, thus reducing the distance that second plunger 302 must be depressed. Sharp end 125 and/or tip 130 should be adapted for tissue penetration as described.

5           Once shaft 120 advances through the abdominal wall, the surgeon grasps it with a hemostat. Pressure is removed from the abdominal wall, delivery device 1 is removed, and anchor 20 is fully deployed. Shaft 120 is exposed above skin surface 601 and barb 102 is set within or on the rectus fascia 603 or other appropriate tissue for later adjustment.

10           The surgeon may now pull (or push) on shaft 120 to raise (or lower) the attached sling until the sling is properly positioned under the bladder neck with suitable tension. Upward movement of shaft 120 may be difficult due to resistance caused by the interleaved first and second ratcheting devices, 140 and 150 (interleaving members 303, 304). When resistance is a problem, the surgeon may position adjuster cylinder 90 around shaft 120, push adjuster cylinder 90 down over shaft 120 through skin surface 601 until adjuster cylinder 90 contacts and  
15           disengages first and second ratcheting devices 140, 150 (interleaving members 303, 304) located within body 99. Adjuster cylinder 90 eases resistance. When the sling is properly tensioned, adjuster cylinder 90 is removed, allowing the first and second ratcheting devices 140, 150 (interleaving members 303, 304) to interleave, resisting movement of shaft 120 with respect to  
20           body 99 and setting the sling tension at this point. Note that it is desirable for several inches of tip 130 to protrude from the abdominal wall to assist in later adjustments. Tension adjustments may be gauged by having indicia markings 80 placed on shaft 120 (see Figures 16a-16h and Figure 18b).

          The exposed end of shaft 120 is taped against the abdominal wall. Postoperatively, the tension on the sling could be adjusted through the operation of the interleaved ratcheting devices  
25           140, 150 (or interleaving members 303, 304) by adjusting the shaft 120 of the tissue anchor protruding through the abdominal skin incrementally, as described above. The patient could be brought back one to several weeks later, and the sling could be adjusted at that time in order to achieve appropriate tension minimizing urethral obstruction postoperatively. The tension is adjusted by axially pulling up or pushing down on shaft 120. Again, if too much resistance is  
30           encountered because of the interleaved ratcheting devices 140, 150 (or interleaving members 303, 304), the resistance can be eased by using adjuster cylinder 90 as described above. The tension

is adjusted to the desired level, and adjuster cylinder 90 is removed, allowing ratcheting devices 140, 150 (or interleaving members 303, 304) to again interleave, resisting movement of shaft 120 relative to body 99. Anchor 20 is now once again substantially fixed in position.

5 The patient's progress is followed, and when a surgeon believes that appropriate tension has been placed, shaft 120 is cut below skin surface 601. This is done with a hollow needle (e.g. 21 gauge beveled) which slides down shaft 120 and cuts shaft 120 in the subcutaneous tissue at the junction with body 99, leaving anchor 20 behind with the sling at the appropriate tension. Alternatively, the exposed portion of the shaft 120 may be cut below skin surface 601 by depressing the skin around shaft 120 and cutting shaft 120. The skin is then released and recoils  
10 leaving the cut tip remote from skin surface 601.

The ability to adjust the tension postoperatively obviates the need for postoperative long-term catheterization. It also obviates the need for re-operation for obstructive uropathy by adjusting the sling postoperatively when the patient is awake and experiences normal daily activities without tissue inflammation. This gives the surgeon a realistic appraisal of the true  
15 tension that is needed for optimum correction of incontinence.

Referring to Figures 17 and 18c-d, adjustment of tissue or sling tension is accomplished by rotation of anchor shaft 120 clockwise or counterclockwise (for increasing or decreasing tissue tension respectively) along the 120 axial shaft. This moves shaft 120 up or down, through body 99 secured to or within tissue by the action of corresponding threads 801 on shaft 120 and  
20 threads 802 on the interior walls of body 99. After appropriate tensioning adjustments were made to shaft 120 trailing through the skin or tissue could be trimmed as previously described with the ratcheted anchor 20. Figure 17 illustrates a similar anchor 20 to the one used in Figures 16a-16g, except the interleaving members on this embodiment include interleaving threads 801, 802. When using interleaving threads 801, 802, however, it is desirable that shaft 120 freely rotate  
25 while attachment member 23 does not. For this reason, attachment member 23 connects to shaft 120 using a ball and socket joint 15.

Figures 16a-16g illustrate a series of drawings depicting one particular embodiment of an adjustable tissue anchor 20 being positioned within a tissue. Figure 16a illustrates anchor 20 placed against the underside of the rectus fascia 603. Delivery device 1 includes shaft 120 having  
30 first interleaving members 303 (or ratcheting devices 150) on its exterior and an attachment member 23. Shaft 120 is positioned within adjuster cylinder 90. Adjuster cylinder 90 and shaft

120 are positioned within hollow chamber 100. Hollow chamber 100 has second interleaving members 304 (or ratcheting devices 140) positioned on its interior walls 121 for interleaving with first interleaving members 303 on shaft 120. Adjuster cylinder 90 keeps first and second interleaving members 303, 304 from engaging. Barb 102 is shown as a set of deployable wings.  
5 Surrounding barbs 102 is a sleeve 96, which is slidably positioned over barbs 102 to keep barbs 102 from deploying before they insert into a tissue.

Viewing Figure 16b, anchor 20 begins to penetrate the rectus fascia 603. The first penetration may occur by cutting edge 90b of adjuster cylinder 90, by sharp end 125 of anchor 20, or by both as shown. Viewing Figure 16c, barb 102 penetrates the rectus fascia 603, slides  
10 through sleeve 96, which remains below the rectus fascia 603. Barb 102 advances into the subcutaneous tissue and deploys.

Viewing Figure 16d, the surgeon pushes adjuster cylinder 90 or anchor shaft 120 to advance cylinder 90 against and through skin surface 601. Alternatively, cylinder 90 may be pressed against the underside of skin surface 601, and the surgeon then accesses cylinder 90  
15 using an exterior incision on skin surface 601. Alternatively, tip 130 may penetrate skin surface 601. Viewing Figure 16e, shaft 120 extends through skin surface 601 where shaft 120 has been pushed through the protruding adjuster cylinder 90 while in the position shown in Figure 16d.

Viewing Figure 16f, the adjuster cylinder 90 has been removed by a surgeon by withdrawing it away from skin surface 601. A button 62 has been positioned relative to shaft 120  
20 and interleaving members 401 on shaft 120 engage interleaving members 402 in center hole 404 of button 62. The interleaving members 401, 402 affix button 62 to shaft 120. As now depicted, anchor 20 is fully set, and barbs 102 resist downward displacement. Upward displacement is resisted by button 62. Shaft 120, however, protrudes through skin surface 601 and may be pulled to increase tension, or pushed to release tension on any device attached to attachment member  
25 23.

Figure 16g illustrates an adjuster cylinder 90 having a funneled upper end 403. Cylinder 90 is reinserted through skin surface 601 to allow shaft 120 to be easily moved without resistance from interleaving members 303, 304. The use of the adjuster cylinder 90 in positioning of shaft 120 is optional.

30 Finally, shown in Figure 20 is a long anchor shaft 120 deployed within a flexible delivery system, such as an endoscope. Generally, anchor 20 will be placed against the desired placement

location, and advanced through the scope, using a rigid or flexible plunger 5 (not shown). Also shown are the video camera 65 and a screen for remote viewing of the procedure.

5 The terms ratcheting devices and interleaving members are used coextensively throughout. It should be noted that interleaving members may be substituted for ratcheting devices and vice versa, depending on a surgeon's desires. As used in the claims, interleaving members is intended to include ratcheting devices. In either case, ratcheting devices or interleaving members include elements that will substantially fix the position of anchor 20 or of shaft 120 positioned within anchor 20. The ratcheting devices and/or interleaving members also allow movement of anchor 20 or shaft 120 so that tension on the sling can be increased by moving shaft 120 or anchor 20 upward or decreased by moving shaft 120 or anchor 20 downward. Ratcheting devices do not exclude elements or configurations that allow shaft 120 or anchor 20 to move in multiple directions.

10 Although the preferred embodiment has been described, it will be appreciated by those skilled in the art to which the present invention pertains that modifications, changes, and improvements may be made without departing from the spirit of the invention defined by the claims.

**CLAIMS**

1. A tissue anchor system comprising a housing, an anchor stay positioned on said housing for retaining an anchor, and an anchor having a barb end with a barb positioned thereon, and an attachment member positioned on said anchor distal from said barb end, said anchor being engageable with said anchor stay to retain said anchor on said anchor stay when said anchor is inserted into a tissue, said barb adapted to resist removal from a tissue once inserted.

2. A tissue anchor system according to claim 1 wherein said anchor further has a shaft connected to said barb end, and said attachment member is positioned on said shaft.

3. A tissue anchor system comprising a delivery device and a tissue anchor, said delivery device having a housing, a plunger slidably positioned in said housing, said anchor having a barb end and a barb positioned thereon, and an attachment member, said anchor being advanced away from said housing upon operation of said plunger, said barb adapted to resist removal from a tissue once inserted.

4. A tissue anchor system according to claim 3 where said anchor further has a shaft attached to said barb end, and said attachment member being positioned on said shaft distal from said barb end.

5. A tissue anchor system according to claim 3 wherein said housing further has a hollow barrel portion, said plunger slidable in said barrel portion, said barb end being sized to be substantially contained within said barrel portion before deployment into a tissue.

6. A tissue anchor system according to claim 3 wherein said shaft is hollow and said plunger has a portion positioned within said hollow of said shaft.

7. A tissue anchor system according to claim 4 wherein said hollow barrel portion of said housing has an axial slot.

8. A tissue anchor system according to claim 6 wherein said shaft of said anchor partially extends through said axial slot in said housing.

9. A tissue anchor system according to claim 3 wherein said barb end has a tip, said tip being shaped to allow said tip to penetrate a tissue.

10. A tissue anchor system according to claim 4 wherein said hollow barrel portion has a tip end, said tip end being shaped to allow said tip end to penetrate a tissue.

11. A tissue anchor system according to claim 3 wherein said housing has a finger grip.

12. A tissue anchor comprising an anchor, said anchor having a barb end and a tip end, a barb

positioned on said tip end, and an attachment member positioned on said distal end.

13. A tissue anchor as in claim 12 where said attachment member comprises a series of projections extending radially from said attachment member.

14. A tissue anchor as in claim 12 where said attachment member comprises a clasp.

5 15. A tissue anchor as in claim 12 where said attachment member comprises at least one opening through said tip end of said anchor.

16. A tissue anchor as in claim 13 further having a button, said button having a center opening adapted to receive said anchor, said center opening of said button having a first interleaving member, said exterior of said anchor having a second interleaving member, said first  
10 and said second interleaving members co-operating when said button is positioned on said shaft to fix the position of said button with respect to said anchor.

17. A tissue anchor as in claim 12 wherein said barb comprises an collapsible umbrella shaped tip.

18. A tissue anchor as in claim 12 further having a retaining device positioned on said barb  
15 end of said anchor.

19. A tissue anchor as in claim 12 where said anchor barb end is positional adjustable with respect to said attachment member.

20. An adjustable tissue anchor comprising a housing, said housing having a barb end and a remote end, said housing having an interior having a first interleaving member, and an anchor shaft, said anchor shaft sized to be positioned in said housing, said anchor shaft having a second  
20 interleaving member, said first interleaving member and said second interleaving member cooperating, when said anchor shaft is positioned in said interior, to adjustably fix the position of said anchor shaft with respect to said housing, said anchor further having an attachment member.

25 21. An adjustable tissue anchor as in claim 20 wherein anchor shaft further has an attachment member.

22. An adjustable tissue anchor as in claim 20 where either said barb end of said housing or said anchor is adapted to penetrate a tissue.

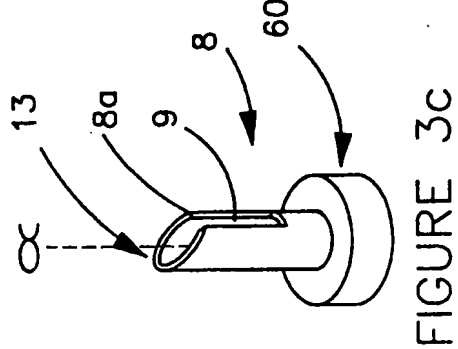
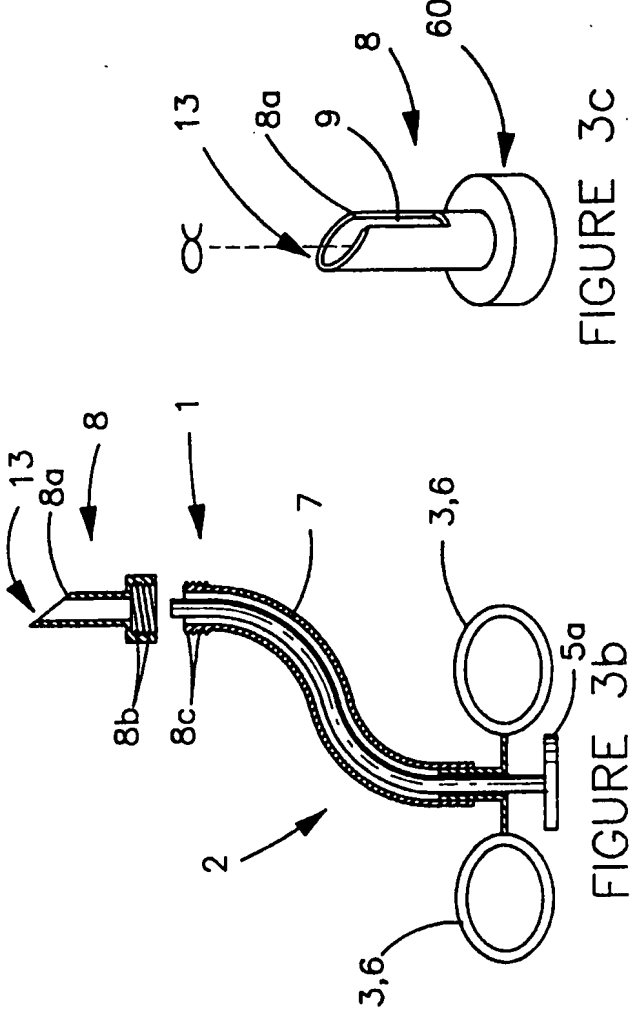
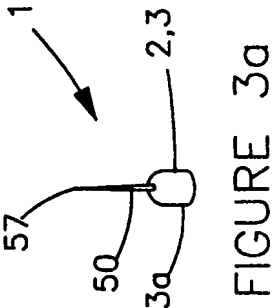
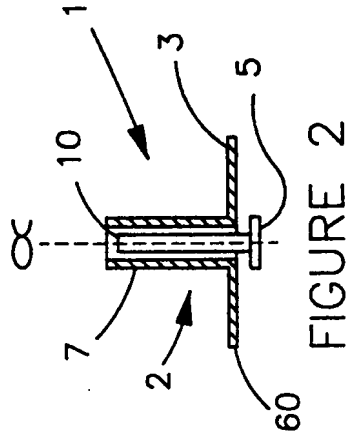
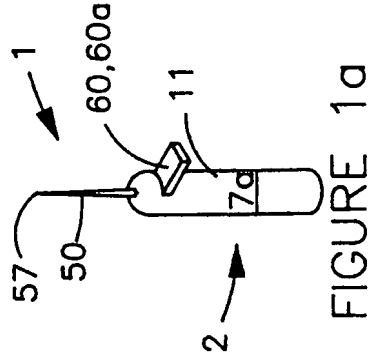
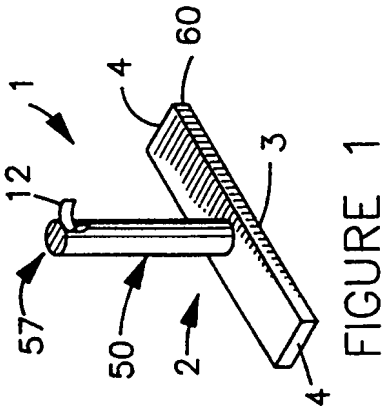
23. An adjustable tissue anchor as in claim 20 where said housing further has a barb  
30 positioned on said barb end.

24. An adjustable tissue anchor system comprising a delivery device and an anchor, said

delivery device having a housing with a hollow barrel portion, said hollow barrel portion having a tip end, said anchor positioned in said hollow barrel portion, said anchor having a barb end and a shaft, a barb positioned on said barb end, said anchor having a hollow chamber axially extending therethrough, said hollow chamber having a first interleaving member, said anchor  
5 further having a central shaft, said central shaft substantially positioned in said hollow barrel, said central shaft being moveably positioned through said hollow chamber of said anchor, said central shaft having a tip end and an attachment end, said central shaft having a second interleaving member, said first and second interleaving members being engageable to resist movement of said central shaft with respect to said anchor.

10 25. An adjustable tissue anchor system according to claim 24 further having a plunger slidably positioned on said housing, said plunger adapted to engage said central shaft when plunged.

26. An adjustable tissue anchor system comprising a delivery device and an anchor, said delivery device having a housing, said housing being adapted to retain said anchor when inserting  
15 said anchor into a tissue, said anchor having a hollow chamber axially extending therethrough, said hollow chamber having a barb end and a shaft, a barb positioned on said barb end, said anchor having a hollow chamber axially extending therethrough, said hollow chamber having a first interleaving member, said anchor further having a central shaft, said central shaft substantially positioned in said hollow barrel, said central shaft being moveably positioned  
20 through said hollow chamber of said anchor, said central shaft having a tip end and an attachment end, said central shaft having a second interleaving member, said first and second interleaving members being engageable to resist movement of said central shaft with respect to said anchor.





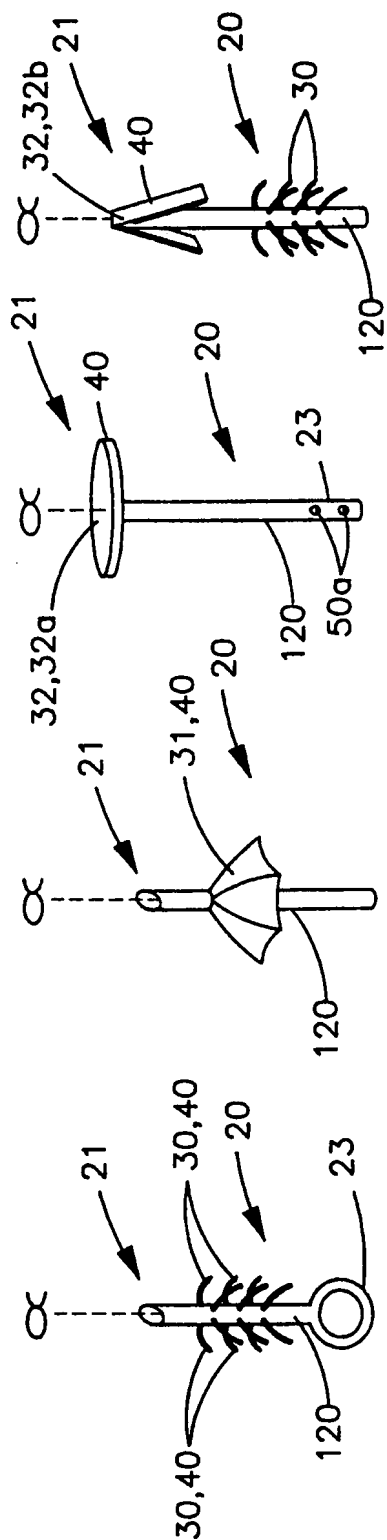


FIGURE 4a

FIGURE 4b

FIGURE 4c

FIGURE 4d

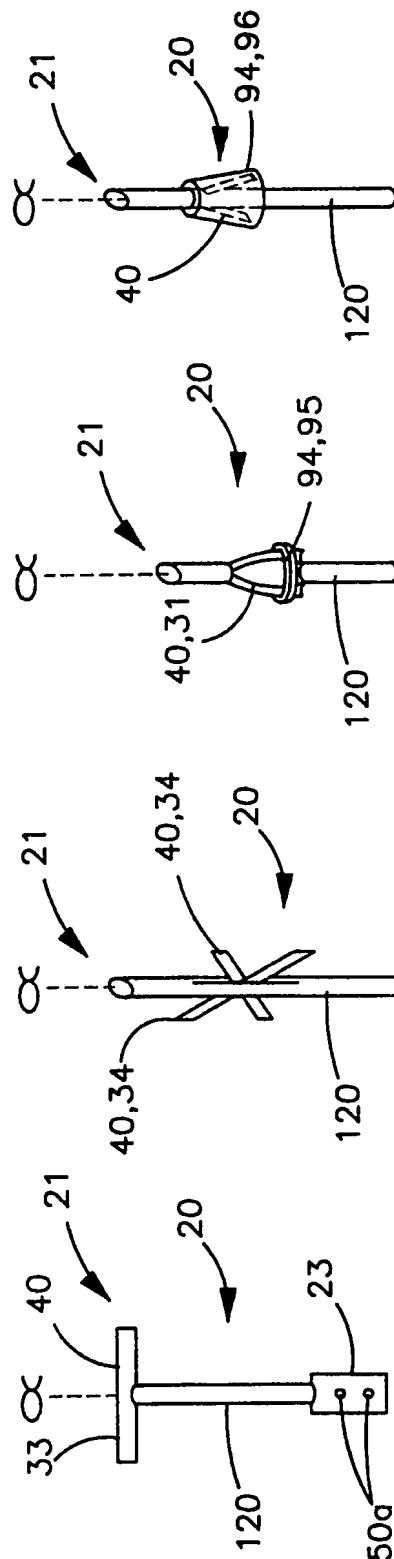


FIGURE 4e

FIGURE 4f

FIGURE 4g

FIGURE 4h

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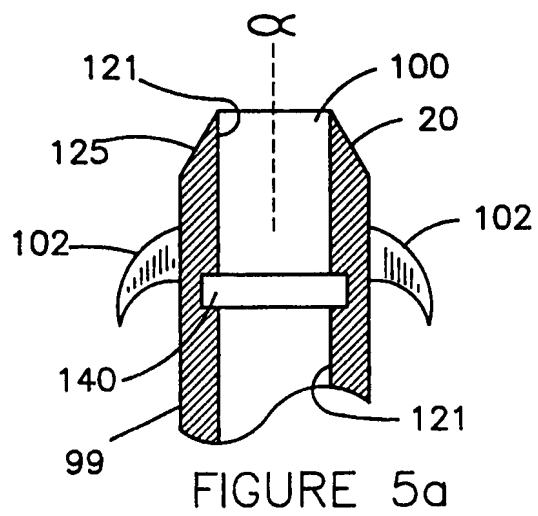


FIGURE 5a

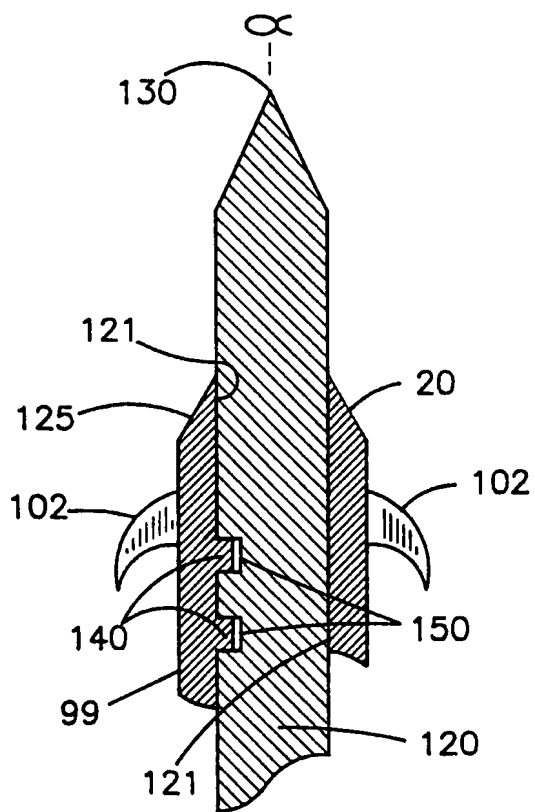


FIGURE 5b

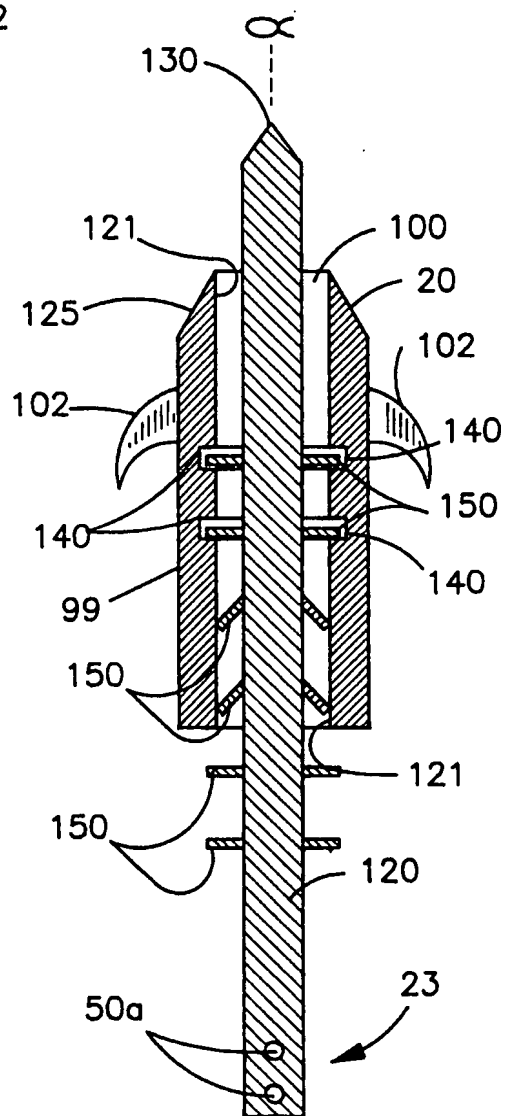


FIGURE 5c

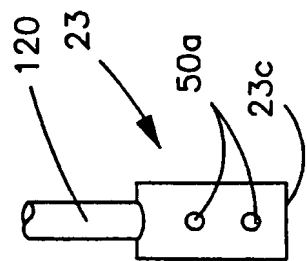


FIGURE 6b

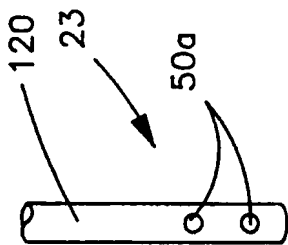


FIGURE 6a

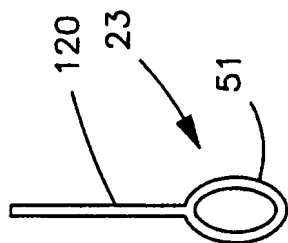


FIGURE 6c

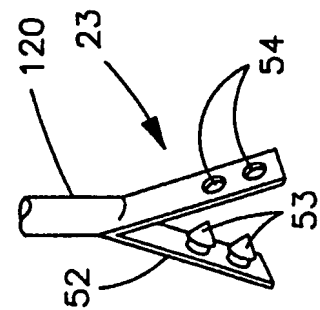


FIGURE 6d

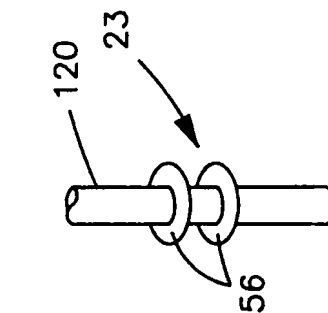


FIGURE 6f

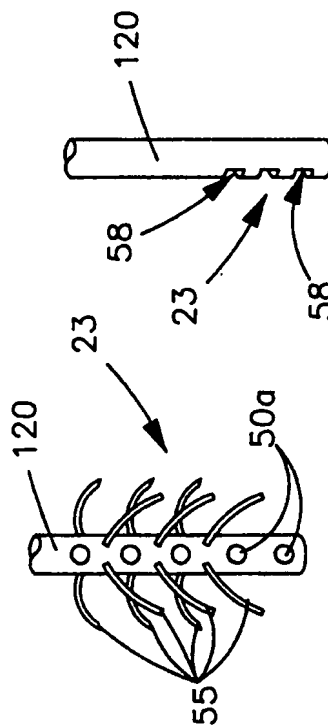


FIGURE 6g

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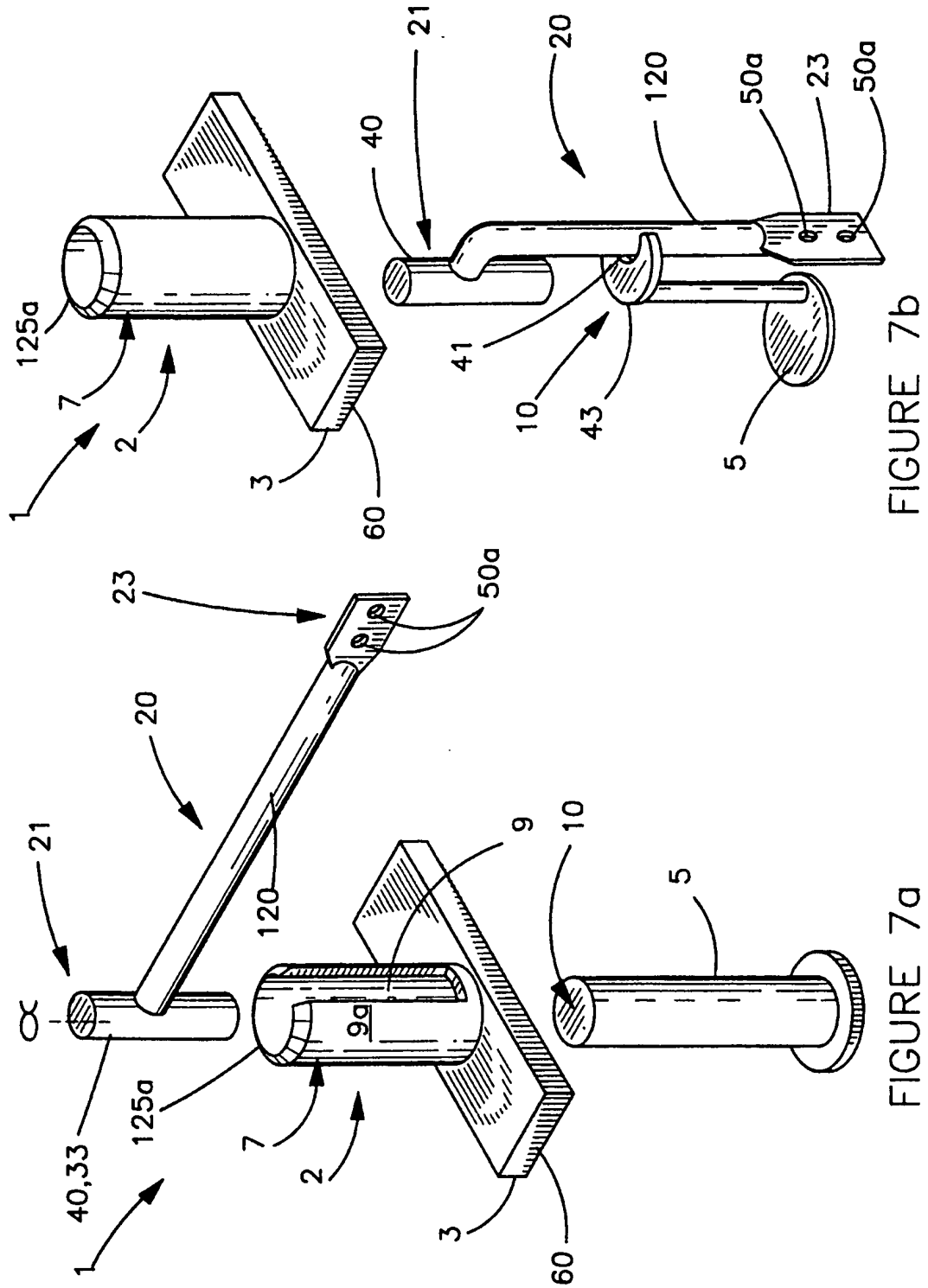


FIGURE 7b

FIGURE 7a

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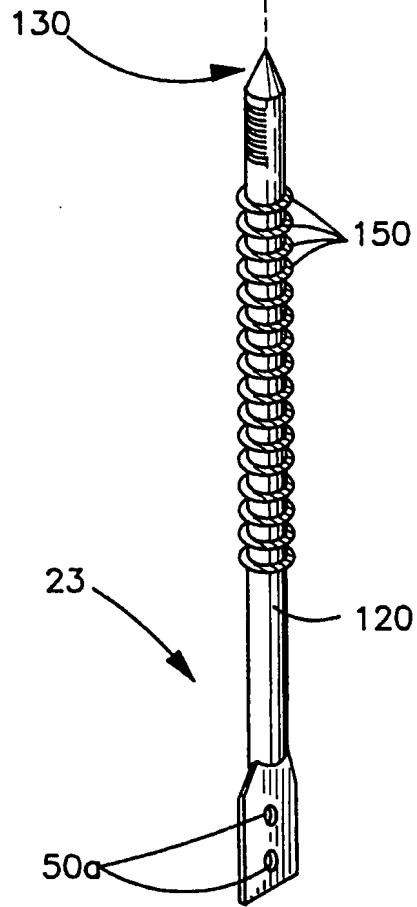
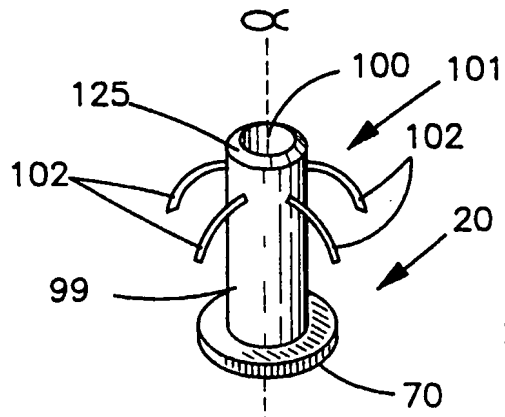


FIGURE 8a

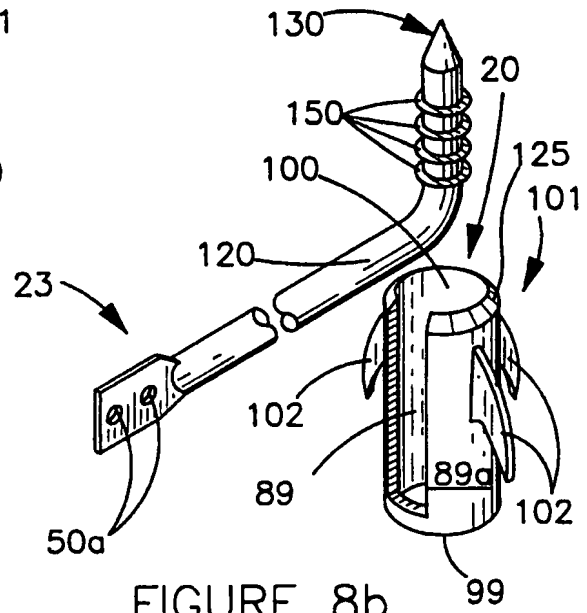


FIGURE 8b

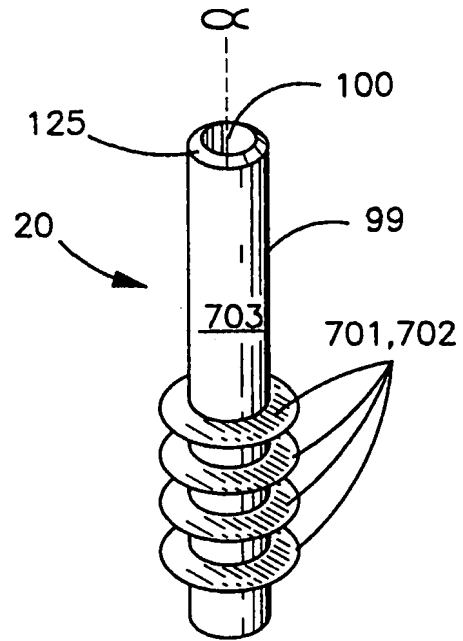


FIGURE 8c

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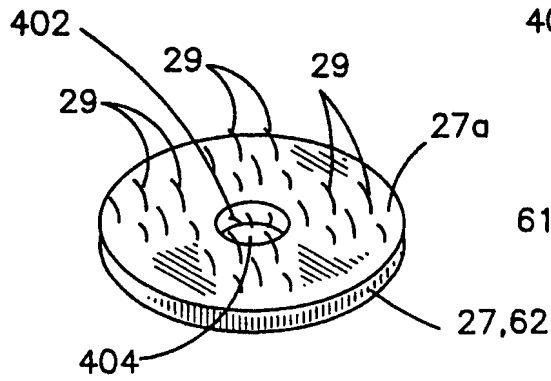


FIGURE 9a

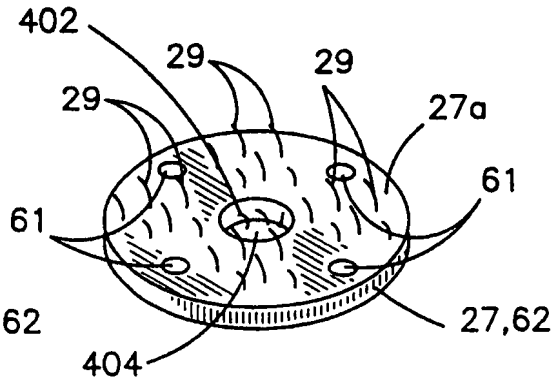


FIGURE 9b

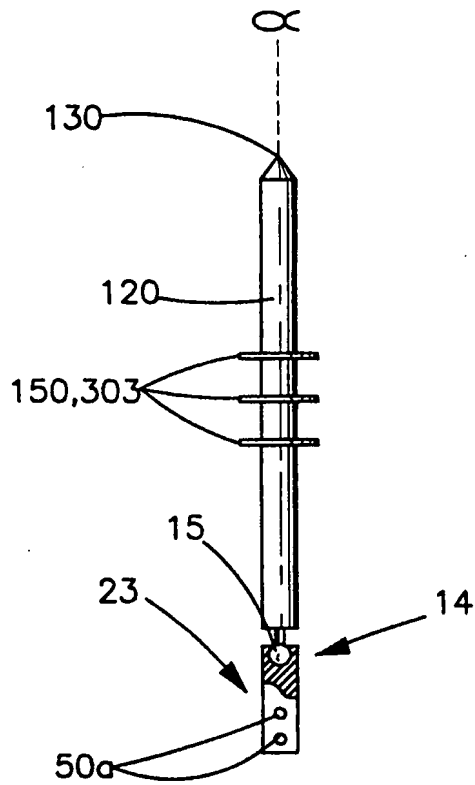


FIGURE 10a

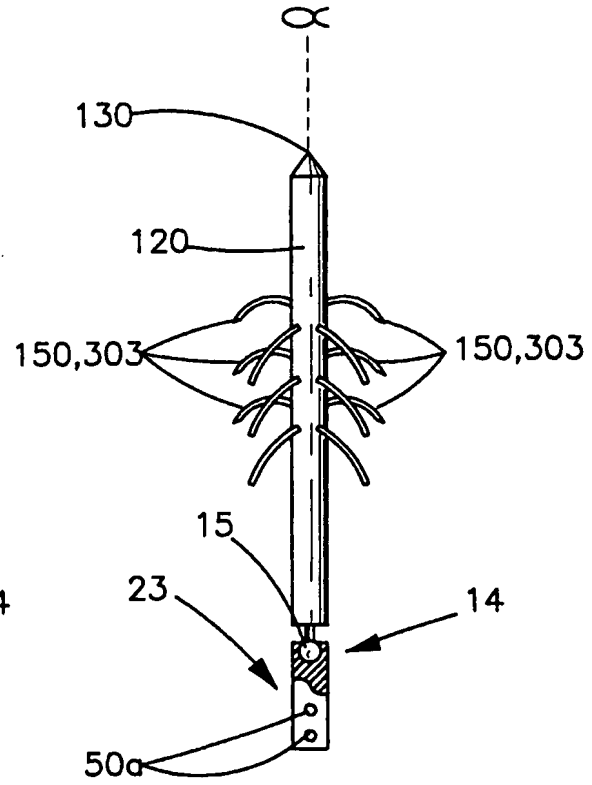


FIGURE 10b

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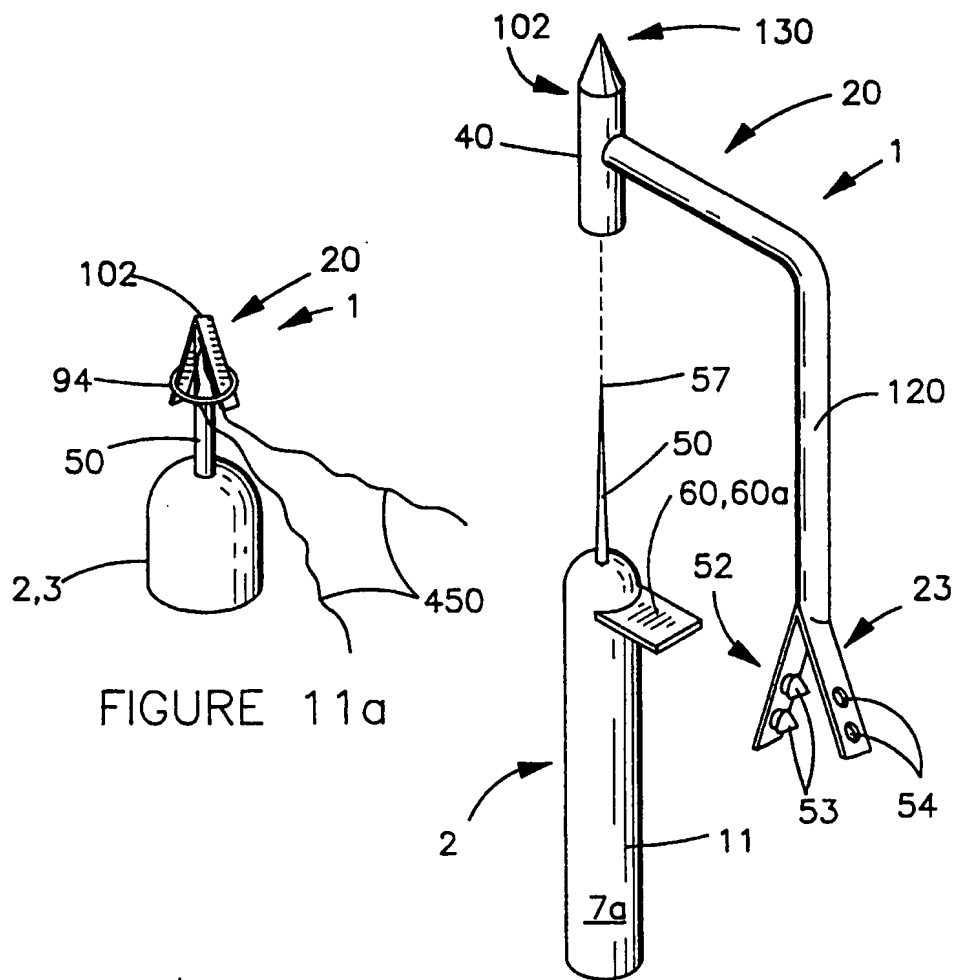


FIGURE 11a

FIGURE 11c

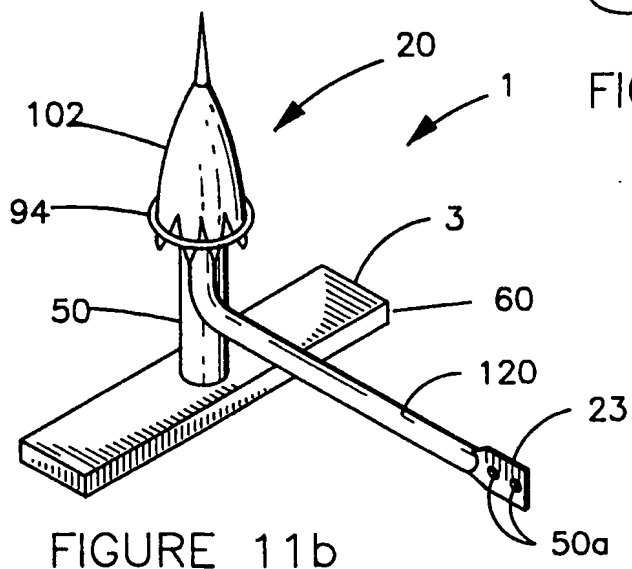


FIGURE 11b

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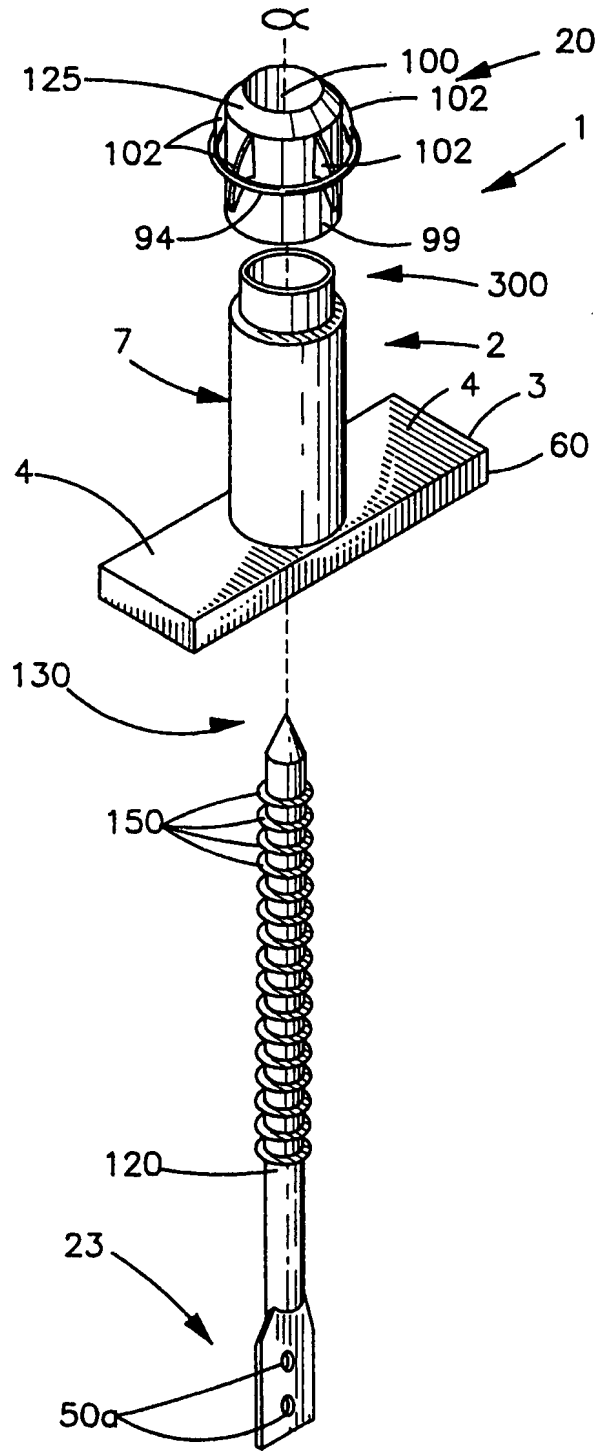


FIGURE 12

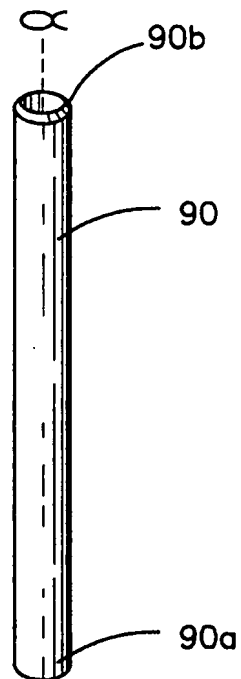


FIGURE 14



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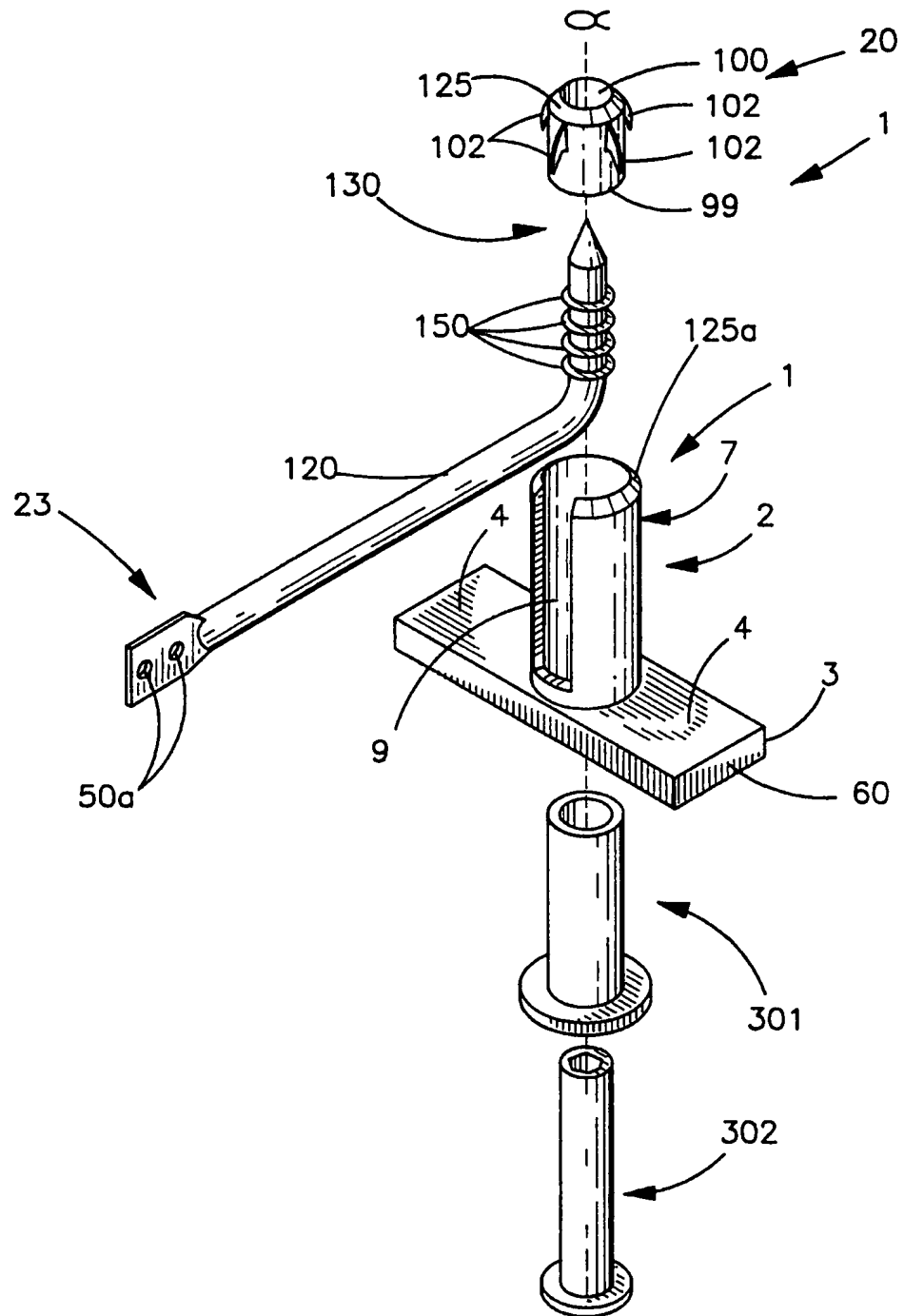


FIGURE 13

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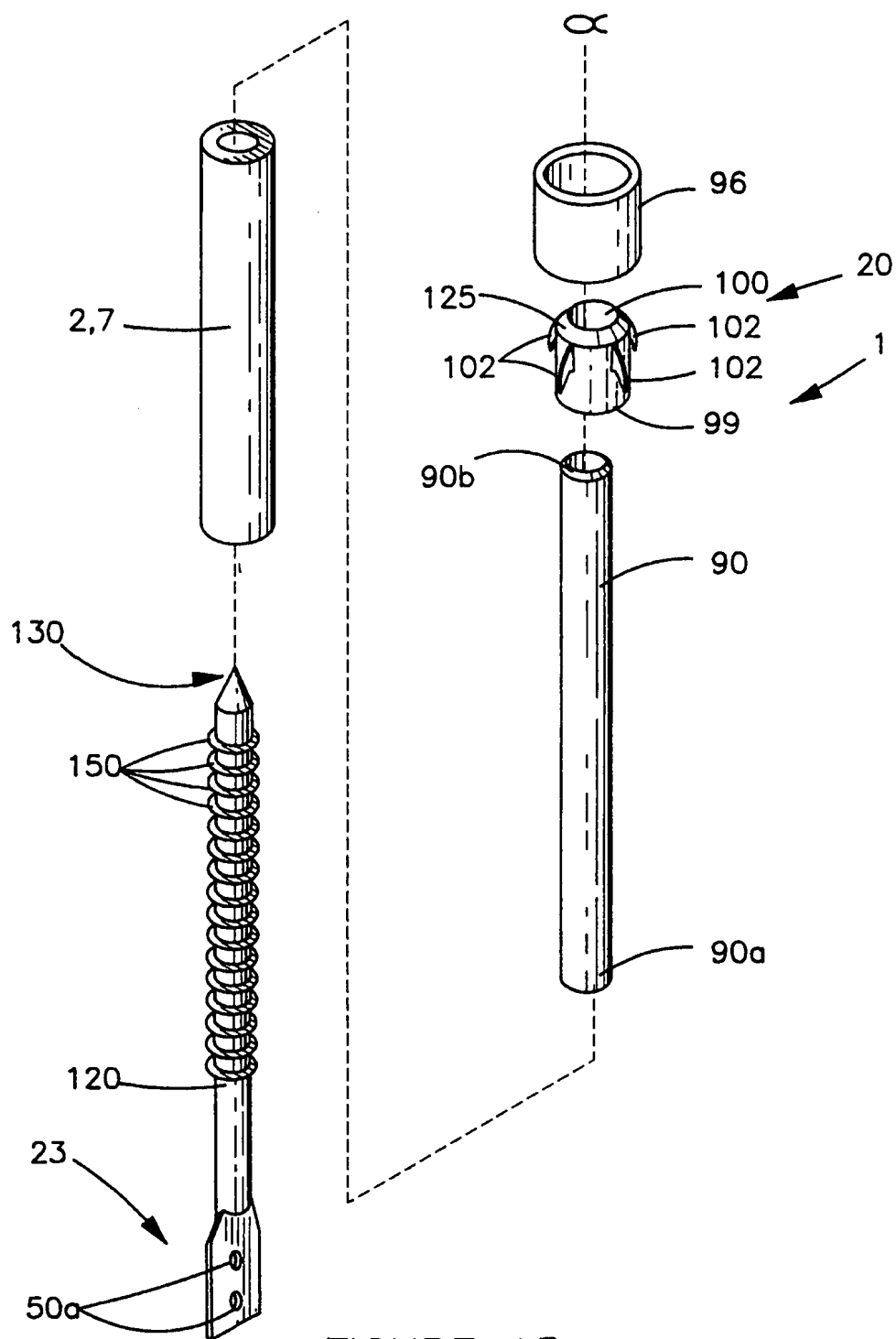


FIGURE 15

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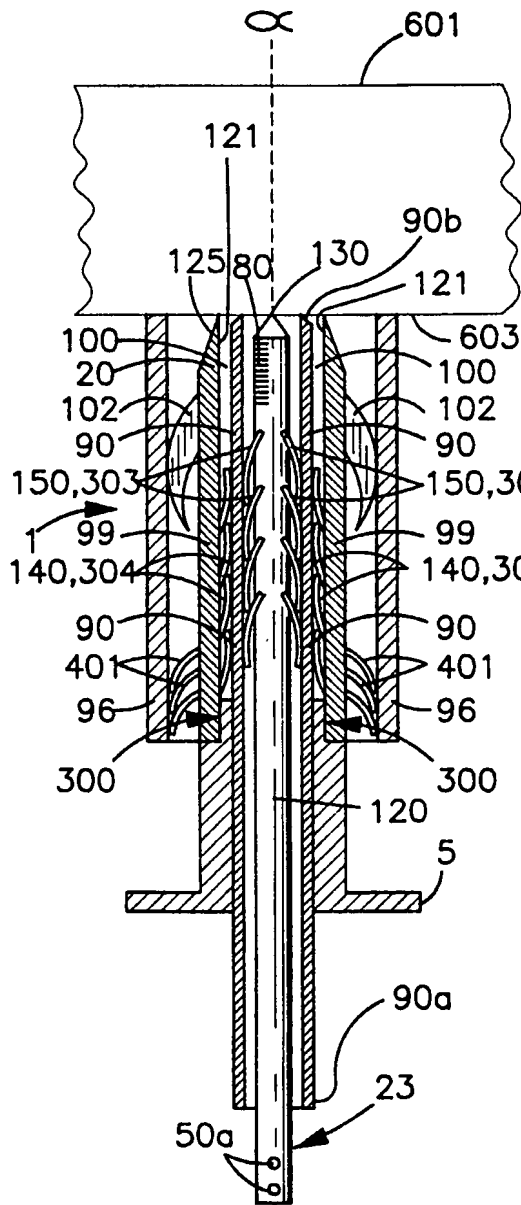


FIGURE 16a

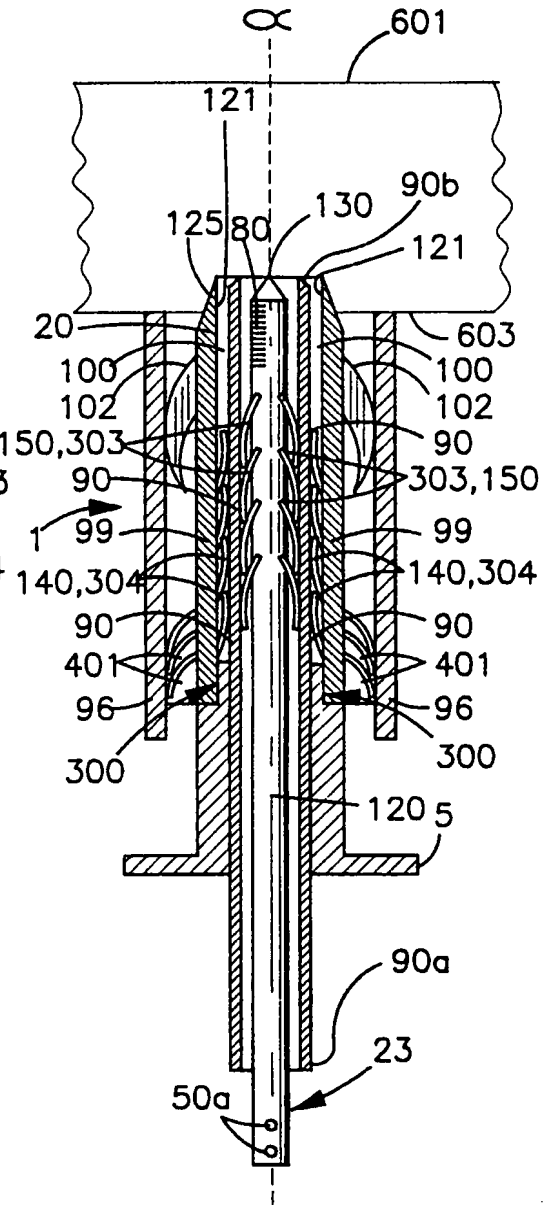


FIGURE 16b

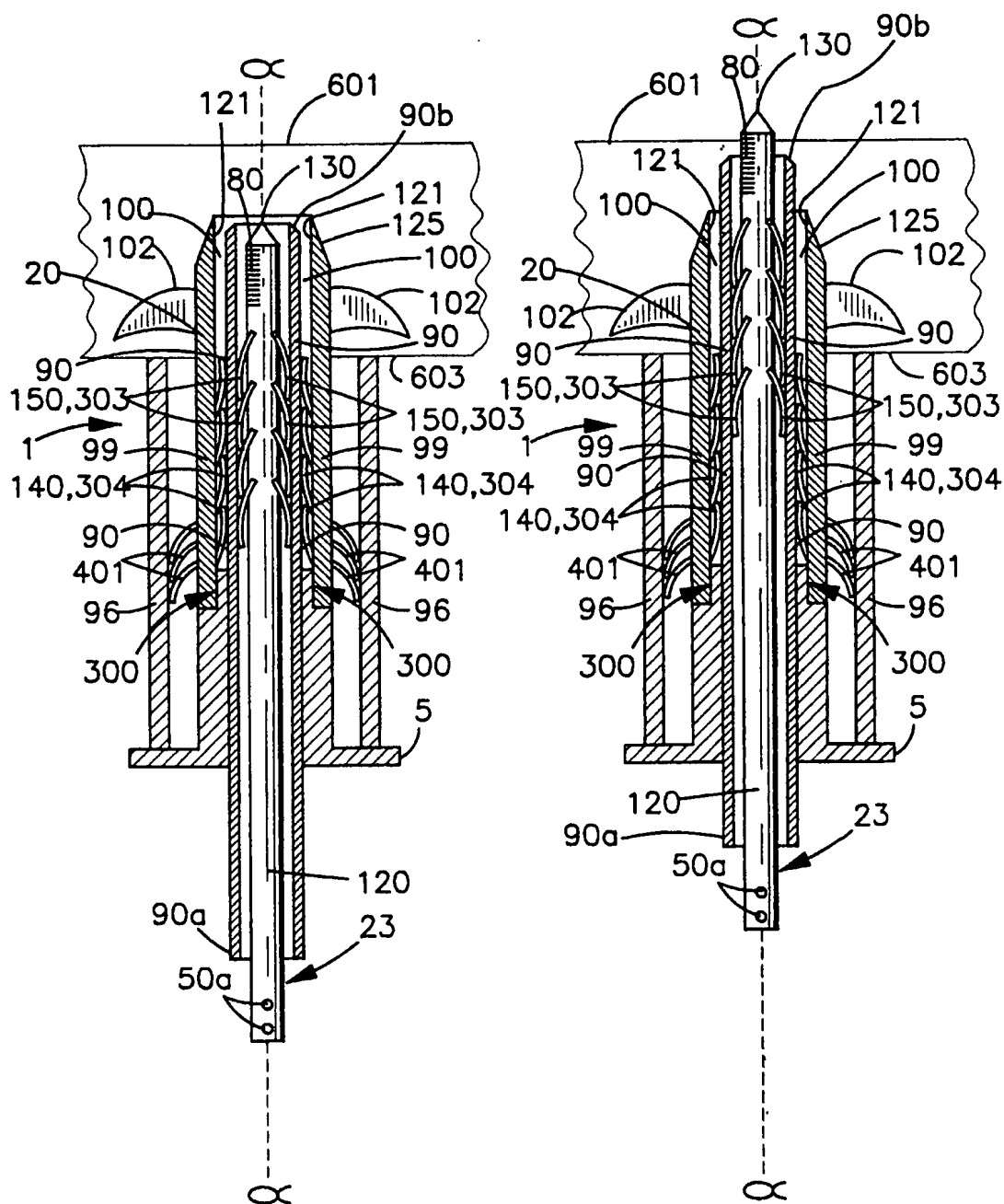
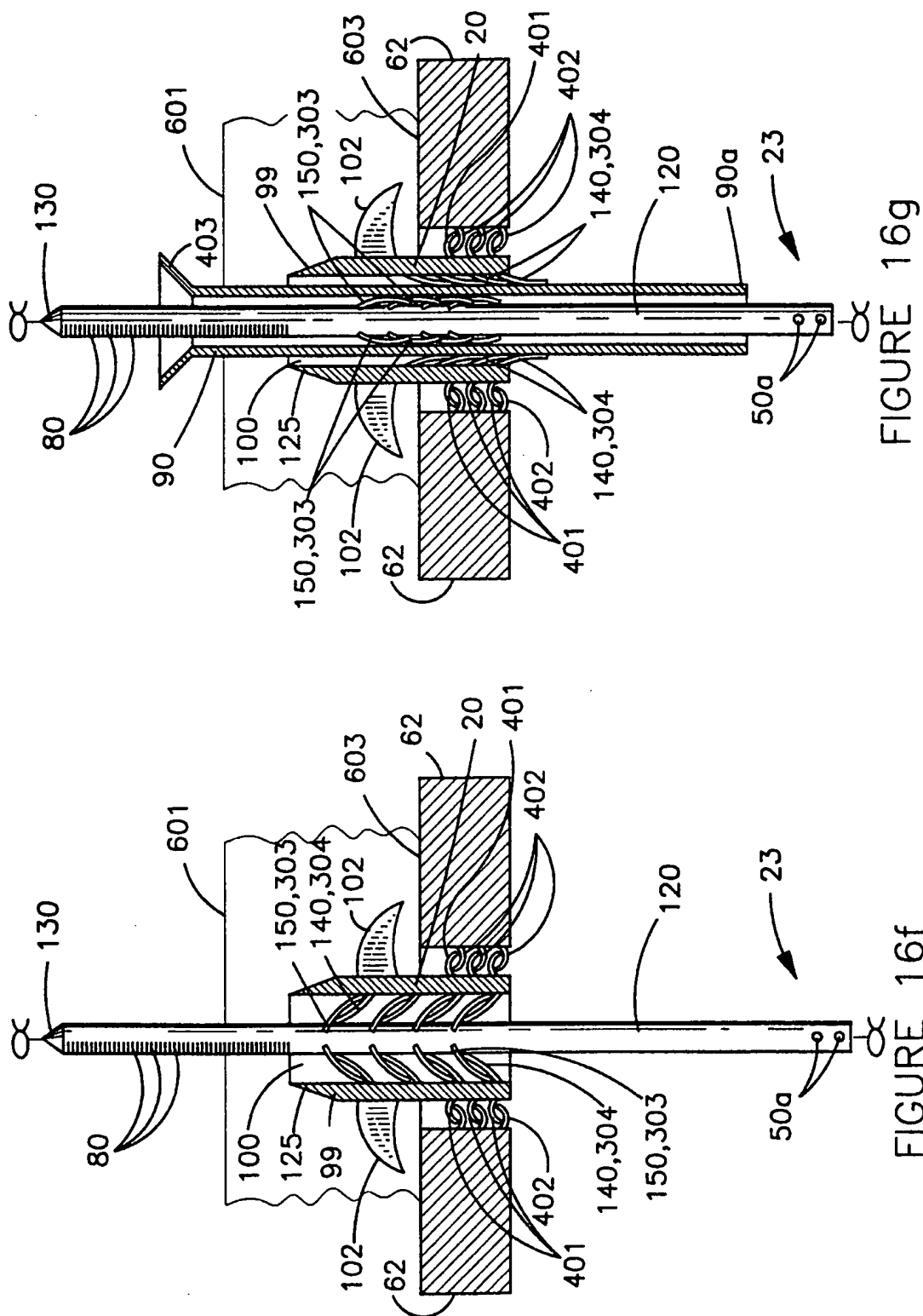


FIGURE 16c

FIGURE 16d





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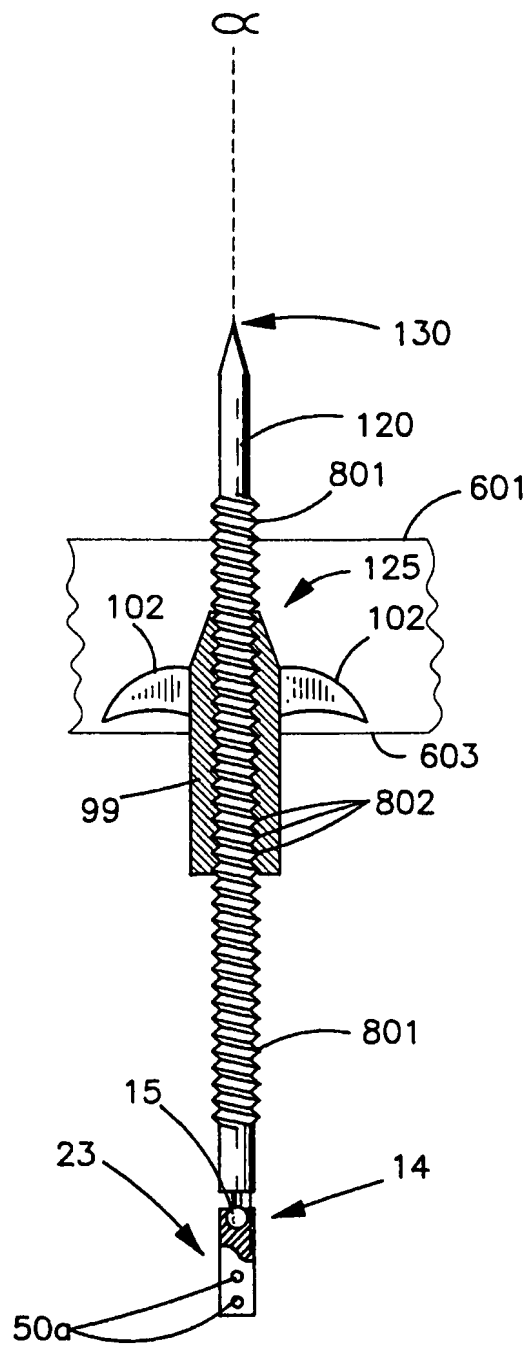


FIGURE 17

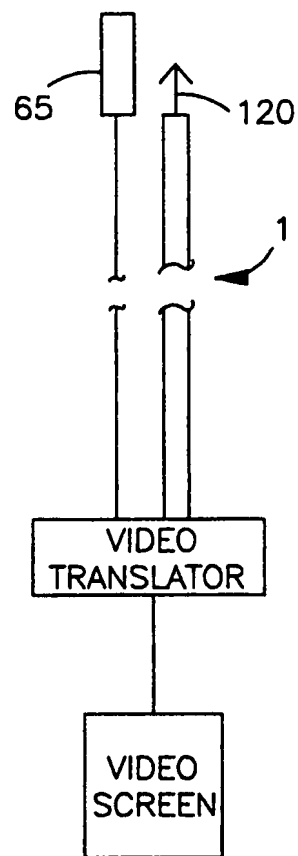
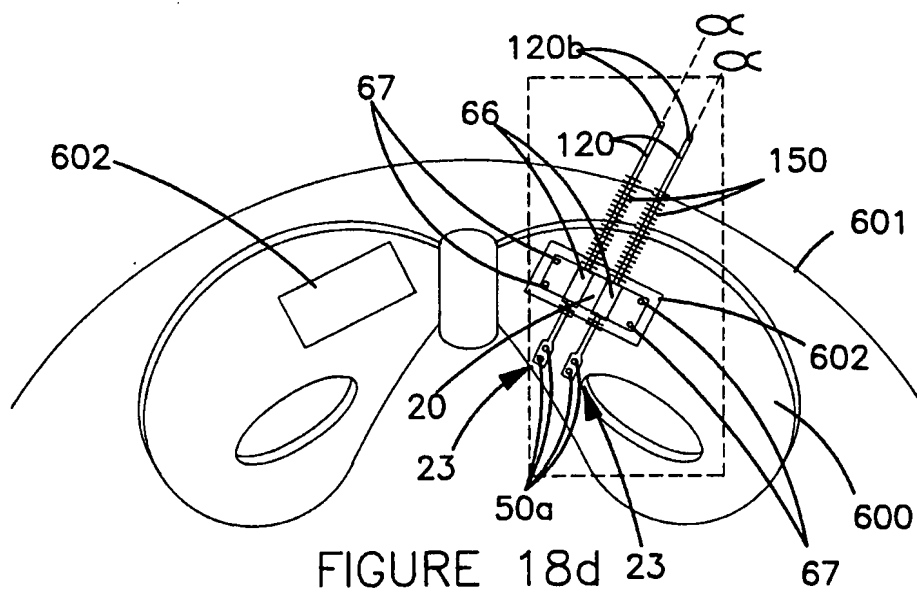
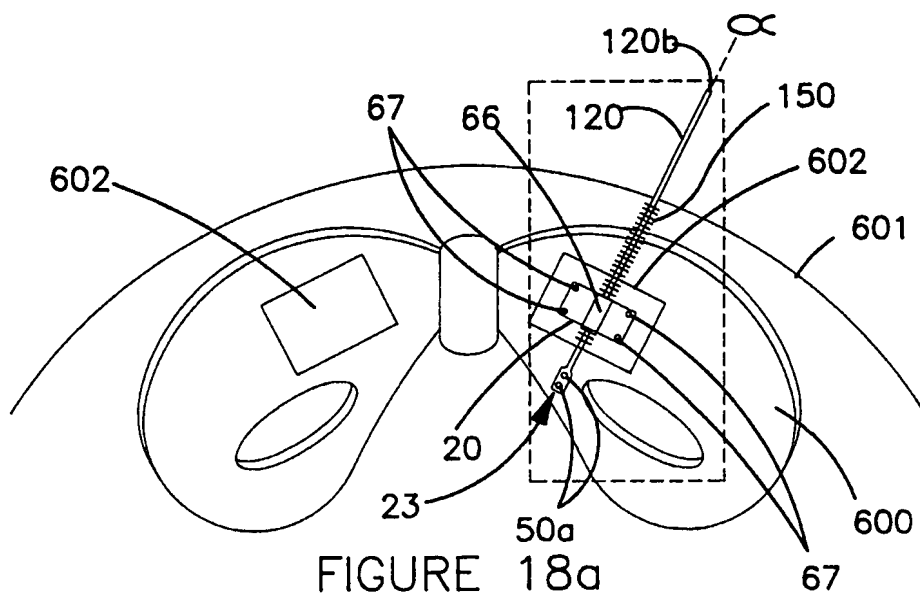


FIGURE 20

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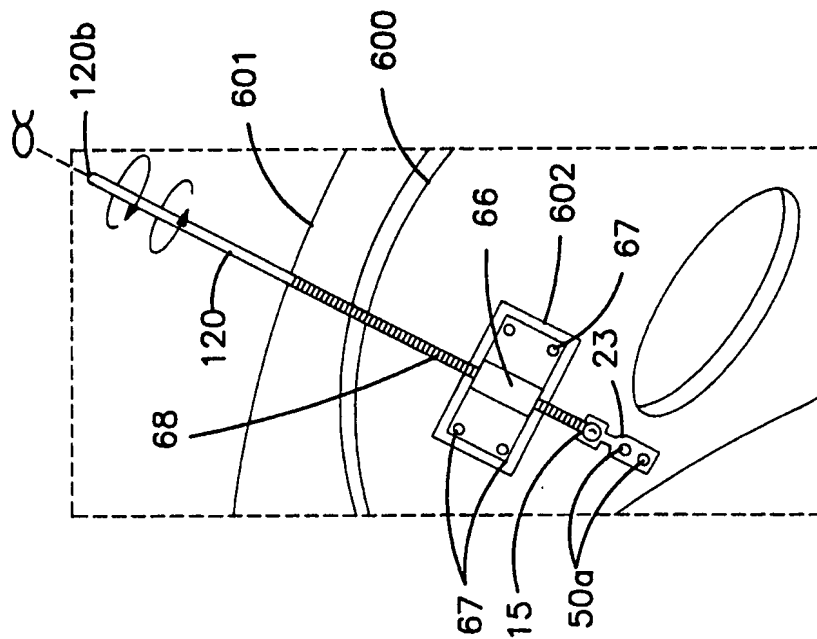


FIGURE 18c

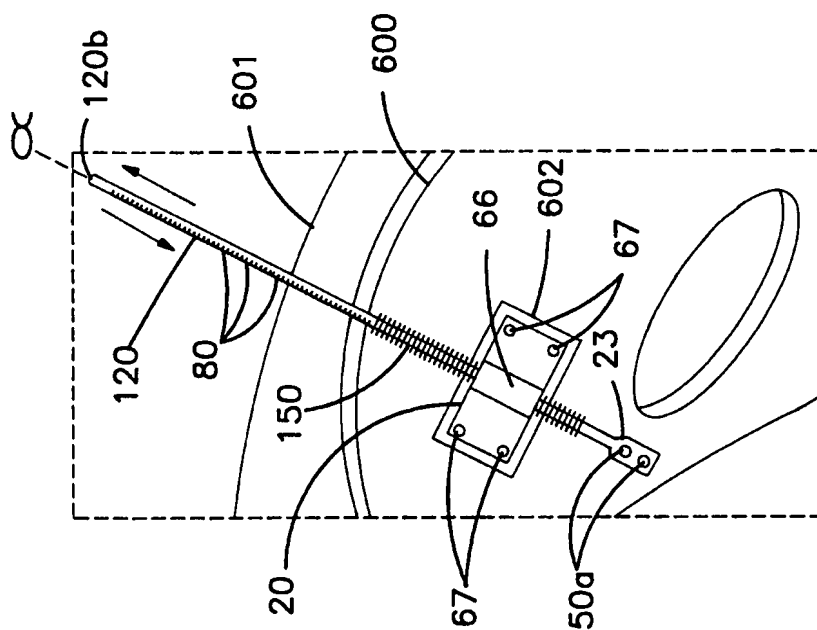


FIGURE 18b

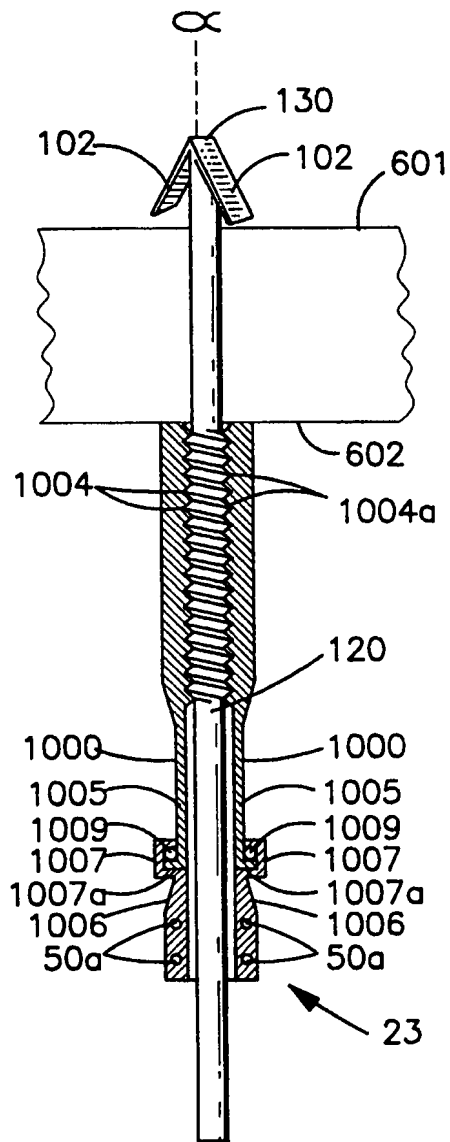


FIGURE 19a

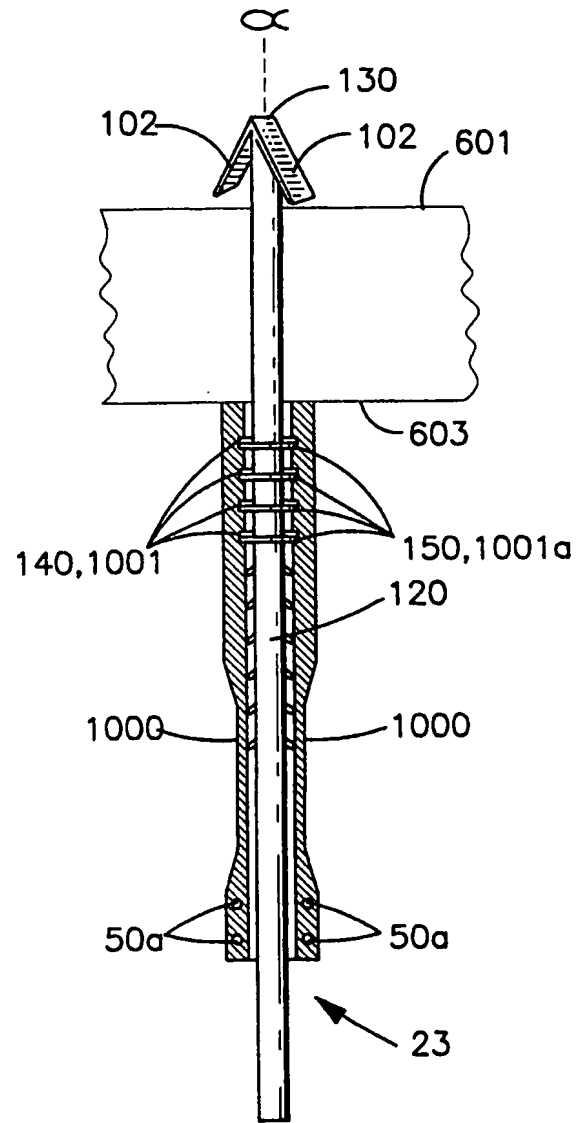


FIGURE 19b

# INTERNATIONAL SEARCH REPORT

International application No.

PCT/US99/11225

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61B 17/04

US CL : 606/72, 232

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 606/72, 232

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y, P	US 5,843,127 A (LI) 01 December 1998, Figs. 5 and 6; and col. 8, lines 6-20.	24-26
Y, P	US 5,849,004 A (BRAMLET) 15 December 1998, Figs. 6 and 28-34; col. 5, lines 12-24; col. 10, lines 28-38; and col. 11, lines 26-68.	1-23

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E* earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Z* document member of the same patent family
*O* document referring to an oral disclosure, use, exhibition or other means	
*P* document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

15 JULY 1999

Date of mailing of the international search report

11 AUG 1999

Name and mailing address of the ISA/US  
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Washington, D.C. 20231

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Authorized officer

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